

**SOIL NEMATODE COMMUNITIES AS INFLUENCED BY FERTILITY
MANAGEMENT PRACTICES, GROWTH AND YIELD OF COMMON
BEANS (var. Mwitemania) IN THARAKA NITHI, KENYA**

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**A Thesis Submitted to Graduate School in Partial Fulfillment of the
Requirements for the Award of the Degree of Master of Science in Soil Science of
Chuka University**

**CHUKA UNIVERSITY
OCTOBER, 2025**

DECLARATION AND RECOMMENDATION

Declaration

This thesis is my original work and has not been presented for any award of a diploma or conferment of a degree in any other institution.

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DEDICATION

This thesis is dedicated to my parents, Josphat Muraki Maina and Nancy Wangui Muraki, my husband, Vincent Kiplagat, and my son, Baraka Kiptanui, who supported my academic success with their financial support, encouragement, endless sacrifices, love, and moral guidance. This work is a testament to their assistance; may Almighty God bless them.

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ABSTRACT

Common bean yield is relatively low due to soil infertility, diseases, and pests such as plant-parasitic nematodes (PPNs), as well as unfavorable climatic conditions, among other factors. PPNs have been reported to cause yield losses of up to 60% while free-living nematodes (FLN) contribute positively to soil health. The study's general objective was to evaluate the effect of fertility management practices on soil nematode communities and the yield of common beans in Tharaka Nithi County. The treatments included NPK (23.23.0), *Trichoderma harzianum* (45g/20 litres), farmyard manure (10 ton/ha), and a control (with no amendment), which were laid out in a randomized complete block design (RCBD) and replicated three times. Soil samples were taken from the field before bean sowing, at flowering, and after harvesting common beans in each trial. Nematodes were extracted from 200ml of soil and 5g of roots using Modified Baermann technique. Data were collected on bean growth every two weeks and on yield components. Meanwhile, nematode numbers were counted, and species diversity and evenness were calculated per plot. The data values obtained were subjected to analysis of variance using R version 4.5.0, and significant means were separated using the Least Significant Difference Test at $\alpha = 0.05$. Results showed that in both trials, Farmyard manure increased FLN in soil (941) and showed a significant difference ($p < 0.05$) from all other treatments and compared to the control, which reduced FLN in soil (110). The control treatment had the highest population of PPNs in soil (773) and showed a significant difference ($p < 0.05$) from the other treatments, while *Trichoderma harzianum* reduced the PPN population (123) in the soil. Control treatment increased Shannon diversity (1.68), Simpson's diversity (0.78), and evenness (0.9) of PPNs as compared to *Trichoderma harzianum*, which reduced Shannon diversity (1.03), Simpson's diversity (0.49), and evenness (0.85). *Trichoderma harzianum* increased the Shannon diversity (1.35), Simpson's diversity (0.7), and evenness (0.84) of FLN compared to the control, which reduced the Shannon diversity (0.85), Simpson's diversity (0.54), and evenness (0.77). The NPK (23.23.0) treatment had the highest growth response which recorded the highest values across four intervals of data collection on height (85.80 cm), number of leaves (72.5), branches (23.54), pods (25.34) seeds (169.39), biomass (1211.72 kg/ha), and grain yield (4592.60 kg/ha), and showed significant difference ($p < 0.05$) from other treatments while control had the reduced growth response of height (39.08 cm), number of leaves (19.25), branches (5.75), pods (5.87) seeds (32.67), biomass (193.24 kg/ha), and grain yield (586.42 kg/ha). *Trichoderma harzianum* had the highest number of nodules (50.5) and showed a significant difference from other treatments, while NPK (23.23.0) had the least (2.9). The NPK treatment increased PPNs in roots (46) and showed a significant difference ($p < 0.05$) from other treatments compared to *Trichoderma harzianum*, which reduced PPNs (2.5) in roots. Farmyard manure and *Trichoderma harzianum* emerged as promising strategies for managing PPNs while promoting beneficial nematodes to enhance common bean yields and soil health. Continuous assessment of nematode communities over multiple growing seasons is crucial for understanding the long-term effects of various soil amendments on diversity and crop productivity.

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ACRONYMS AND ABBREVIATIONS

| | |
|----------------|--|
| ANOVA | : Analysis of Variance |
| DAP | : Diammonium Phosphate |
| FAO | : Food and Agriculture Organisation of the United Nations |
| FAOSTAT | : Food and Agriculture Organisation Corporate Statistical Database |
| FLN | : Free-living Nematodes |
| ICIPE | : International Centre of Insect Physiology and Ecology |
| KARLO | : Kenya Agricultural Research and Livestock Organisation |
| MSW | : Municipal Solid Waste |
| NACOSTI | : National Commission for Science, Technology and Innovation |
| NPK | : Nitrogen Phosphorus Potassium |
| PPNs | : Plant-Parasitic Nematodes |
| WUE | : Water-Use-Efficiency |

CHAPTER ONE

INTRODUCTION

1.1 Background Information

Common beans (*Phaseolus vulgaris*) belong to the Fabaceae family, which includes other legumes such as soybeans, peas, chickpeas, and alfalfa. Common beans account for over $\frac{3}{4}$ of the world's annual production of 8.5 million metric tons (Rodriguez & Creamer, 2014). In Africa, Tanzania is the largest producer of common beans, followed by Uganda, Kenya, Ethiopia, and Rwanda, accounting for 17.52, 15.28, 11.16, 8.25, and 6.59% respectively (Mathobo & Mathobo, 2024). In Kenya, common beans are mainly produced in the Western, Eastern, Central, and Rift Valley regions, accounting for 31, 24, 20, and 13%, respectively, on an estimated 700,000 ha (Farrow *et al.*, 2020).

Beans are significant because they provide complex carbohydrates, protein, and micronutrients that are valuable to the human population (FAOSTAT, 2019). Farmers use common beans in crop rotation because nitrogen is an essential nutrient for plant growth. Common beans improve soil fertility through nitrogen fixation, which means they harbor rhizobium bacteria in their nodules that convert atmospheric nitrogen into plant-available forms (FAOSTAT, 2019). Bean yields in Kenya are as low as 750 kg/ha in monocropping and even lower than 335 kg/ha under mixed cropping systems, whereas the potential is 1500-2000 kg/ha (Mataa *et al.*, 2021). This could be attributed to pests like plant-parasitic nematodes, low soil fertility, poor varieties, and water stress, among other agronomic factors. The potential for nodulation of leguminous crops is adversely affected by infection with root-knot nematodes, thereby interfering with nitrogen fixation. According to Talwana *et al.* (2016), common bean yield losses have been recorded by up to 60% due to heavy infestation of root-knot nematodes.

Nematodes are worms that are microscopic and live in soil pores formed through soil processes (Yeates *et al.*, 2009). They move in water films, which stick to soil particles. The estimated population of species is more than half a million, whereas free-living species have the largest populations in soils, freshwater habitats, and oceans. At the same time, parasitic types form the smaller group (Ferris *et al.*, 2012). Soil nematode communities comprise free-living nematodes and plant-parasitic

nematodes. Free-living nematodes are useful indicators of soil health (Yeates, 2003), and entomopathogenic nematodes are utilized for the biological control of insects (Schratzberger *et al.*, 2019; Neher, 2001). They play crucial roles in the soil, like providing food sources to soil microbes, improving water holding capacity and soil structure, mineralization, and decomposition of organic matter (Lu *et al.*, 2020; Nisa *et al.*, 2021). They are primarily found in marine and soil environments. Nematodes are classified into five trophic levels: omnivores, bacterivores, herbivores, fungivores, and predators (Yeates *et al.*, 1993; Zullini, 2018). Plant-parasitic nematodes (PPNs) are considered a primary biological constraint worldwide in various crops, resulting in considerable losses in production of up to 20% (Hamida *et al.*, 2015). A key plant-parasitic nematode, the *Meloidogyne* spp. cause losses of up to 85% (Coyne *et al.*, 2014). *Meloidogyne javanica*, *M. incognita*, and *M. arenaria* are the most encountered in tropical regions and have a wide host range (Bebber *et al.*, 2014).

Strategies like cultural practices, nematicides, organic amendments, and resistant varieties have been established to manage plant-parasitic nematodes (Sikora & Roberts, 2018). Some of these strategies have limitations, such as nematicides, which are costly and not environmentally friendly (Sasaneli *et al.*, 2021). Cultural practices are labor-intensive and time-consuming. Resistant varieties may not be efficient when a field has multiple nematode species (Oka *et al.*, 2000). Organic amendments, such as farmyard manure, suppress plant-parasitic nematodes, increase free-living nematodes, and improve soil health due to the added nutrients in the soil (Thoden *et al.*, 2011). The addition of farmyard manure influences nematodes due to the incorporation of organic matter; nematodes can respond to manure amendments by increasing their activity levels and numbers (Semenov *et al.*, 2021).

Farmyard manure can indirectly affect soil organisms through changes in physical properties (such as porosity and aggregation), soil pH, soil organic matter levels, and productivity (Bunemann *et al.*, 2006). Chemical fertilizers influence soil organisms, stimulating the growth of nitrifying and ammonifying microorganisms, as well as the proliferation of bacteria that form spores and promote crop residue mineralization (Pahalvi *et al.*, 2021). Chemical fertilizers, such as NPK, are added to enhance soil fertility (Bhatt *et al.*, 2019). These fertilizers can influence both parasitic and free-

living nematodes, often resulting in shifts in trophic structure, diversity, and abundance (Fan *et al.*, 2025).

Trichoderma harzianum is a biocontrol agent used for biostimulation and against phytopathogens in agriculture. *Trichoderma harzianum* is used as a seed treatment, foliar application, and soil application for suppressing fungal pathogens (Pani *et al.*, 2021). Applying *Trichoderma harzianum* to the soil has several benefits, including creating a barrier around the roots, improving soil health, and enhancing the availability of nutrients (Eman *et al.*, 2023). *Trichoderma* has been shown to promote plant growth in many ways. It has been used as a biofertilizer because it stimulates the plant growth of various crops. It can be used as an alternative to chemical fertilizers or as an amendment to improve crop production (Akladios & Abbas, 2014). *Trichoderma* strains produce asexual spores only, so they don't have a sexual stage. However, for very few strains, the sexual stage is known, but not in the strains used as biocontrol agents (Ghazanfar *et al.*, 2018). Its efficacy is affected by both abiotic and biotic factors, and its efficacy is maximised by microencapsulation (Maruyama *et al.*, 2020). It produces volatile compounds and plant growth hormones. It solubilizes phosphates that are unavailable to the plant. It promotes the uptake of micro- and macro-nutrients required by the crop (de Lima Gonilha *et al.*, 2024).

Therefore, the study aimed at assessing how different soil amendments influence soil nematode populations as indicators of soil health. It also sought to determine whether these amendments improve the yield of common beans. Soil nematode populations were monitored to understand their role in soil and crop performance under different treatments. This study enhanced the identification and counting of free-living and plant-parasitic nematodes. Therefore, a study was carried out to evaluate the influence of the soil fertility management practices, growth, and yield of common beans on nematode communities in Tharaka Nithi County, Kenya.

1.2 Statement of the Problem

Common bean yield in Kenya is relatively low due to low soil fertility, pests and diseases, and unfavourable climatic conditions, among other factors. The total food demand is expected to double by 2030, and plant-parasitic nematodes have been reported to reduce yields. Plant-parasitic nematodes can result in yield losses of up to

60%. For instance, root-knot nematodes affect the nodulation of common beans. More research has been conducted on the impact of plant-parasitic nematodes on the yields of common beans; however, beneficial nematodes, such as free-living nematodes, have received relatively little attention. Free-living nematodes play a crucial role in the decomposition of organic matter and the recycling of nutrients. For instance, nematode fungivores and bacterivores do not feed directly on soil organic matter, but rather on fungi and bacteria that decompose organic matter. Scarce information is available on the influence of soil fertility management practices on soil nematodes. There is also scarce information on the abundance and diversity of plant-parasitic nematodes and free-living nematodes in Tharaka Nithi County.

1.3 Objectives

1.3.1 General Objective

To evaluate soil nematode communities as influenced by fertility management practices, growth and yield of common beans in Tharaka nithi, County.

1.3.2 Specific Objectives

- i. To determine the effects of NPK (23.23.0), *Trichoderma harzianum*, and farmyard manure on numbers, evenness, and species diversity of soil nematodes under common beans
- ii. To determine the effect of NPK (23.23.0), *Trichoderma harzianum*, and farmyard manure on common beans' growth and yield components.
- iii. To determine the effect of NPK (23.23.0), *Trichoderma harzianum*, and farmyard manure on plant-parasitic nematodes in the roots of common beans.

1.4 Hypotheses

H₀₁: There is no statistically significant effect of NPK (23.23.0), *Trichoderma harzianum*, and farmyard manure on numbers, evenness, and species diversity of soil nematodes under common beans.

H₀₂: There is no statistically significant effect of NPK (23.23.0), *Trichoderma harzianum*, and farmyard manure on the growth and yield components of common beans.

H₀₃: There is no statistically significant effect of NPK (23.23.0), *Trichoderma harzianum*, and farmyard manure on plant-parasitic nematodes in the roots of common beans.

1.5 Justification

Common bean plays a vital role in human nutrition, although its production is affected by plant-parasitic nematodes, resulting in low yields. *Trichoderma harzianum* is a biocontrol agent for plant-parasitic nematodes, a growth-promoting agent in common beans, and a biofertilizer. It is an essential component for managing plant-parasitic nematodes, which are economically important (Maruyama *et al.*, 2020). Free-living nematodes are beneficial and crucial to plant growth by decomposing organic matter and cycling nutrients in soil. They feed on living tissues, algae, fungi, bacteria, and dead organisms (Schratzberger *et al.*, 2019). They improve the water-holding capacity and soil structure and release nutrients to plants. Nematodes serve as biological indicators because their community structure can indicate soil health (Marais *et al.*, 2020). Farmyard manure and NPK (23.23.0) increase soil fertility once applied to the soil and hence will increase common bean yields. Inorganic fertilizers release nitrogen and phosphorus, encouraging nematode multiplication and plant growth (Pahalvi *et al.*, 2021). The addition of farmyard manure leads to checked nematode populations because of competition for space and food with the soil organisms introduced. Due to adding organic matter, nematodes can react with manure amendments by increasing activity levels and numbers (Semenov *et al.*, 2021). There is a need to grow common beans in Tharaka Nithi County for several reasons, including enhancing nutrition and food security, improving the livelihoods and income of farmers, improving soil fertility, and promoting agricultural sustainability (Lisciani *et al.*, 2024). Farmers are advised on the appropriate strategies for managing plant-parasitic nematodes and promoting beneficial ones to enhance the yields of common beans and contribute to food and nutrition security.

CHAPTER TWO

LITERATURE REVIEW

2.1 Overview of Common Beans Production

Common beans, *Phaseolus vulgaris*, were domesticated in South America and Central Mexico 8000 years ago (Gaut, 2014). Ecuador and Peru were the centres of origin of common beans (Morales, 2006). India is the world's largest producer of beans. In 2021/2022, India produced 6,120,000 tonnes of dry common beans (Delfini *et al.*, 2021). Tanzania is Africa's largest producer of common beans, producing 1.2 tons annually (Onyango, 2022). Common beans are widely cultivated in Kenya, especially for urban and rural diets. Common beans are mainly produced in Central, Western, Eastern, and Rift Valley regions (Duku *et al.*, 2020). Despite their caloric intake, common beans are an essential source of protein and micronutrients (Nadeem *et al.*, 2021). Common beans are appreciated widely because they are affordable and have a long storage life.

In Kenya, common beans are mainly intercropped with maize, cassava, coffee, millet, potatoes, and sorghum. The ecological requirements of common beans: soil pH of 6.5-7.5, altitude of 1000-2100m asl, rainfall of 800-2000mm, and temperature of 20-25°C; the soils should be well drained and not waterlogged (Porch *et al.*, 2013). Constraints of common bean production are inadequate seeds, labour, diseases, pests, drought, and animal damage (Mangole *et al.*, 2022). According to Mwaniki (2002), the unavailability of seeds limits the production of common beans. Most farmers use uncertified and of poor quality seeds; hence, there is a need to train farmers on quality seed production. Some pests affecting common beans in Kenya are plant-parasitic nematodes, aphids, armyworms, corn earworms, cutworms, leaf miners, loopers, Mexican bean beetles, stinkbugs, and spider mites (Singh & Schwartz, 2011).

Nematodes are free-living in soil, marine, and freshwater environments, but many species are parasitic, particularly on various animals and plants. The parasitic species have veterinary, clinical, and agricultural importance as parasites of livestock and man and pests of plants (Sánchez-Moreno & Ferris, 2018). Nematodes are found at the bottom of rivers, lakes, and huge depths, especially in oceans. Some nematode species can withstand temperatures consistently below the freezing point, while others thrive in hot springs. Nematode infestations at high initial population density result in

significant yield losses. For instance, according to Adomako *et al.* (2022), yield losses may go up to 10%- 80%, especially with lesion nematodes, as well as 50%- 90% with root-knot nematodes (*Meloidogyne* spp.). Plant-parasitic nematodes, like the root-knot nematodes, predispose various crops to soil-borne microorganisms, which cause wilt diseases and root rot. Root-knot nematode infection also suppresses nodulation (Kimenju *et al.*, 1999). Common beans are susceptible to various diseases caused by fungal oomycetes, bacteria, fungi, and viruses (Gupta & Singh, 2020).

2.2 Effects of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Numbers, Evenness, and Species Diversity of Soil Nematodes under Common Beans

2.2.1 Populations, Evenness, and Diversity of Soil Nematodes under Common Beans

According to Adomako *et al.* (2022), who conducted research to identify the occurrence of nematodes associated with common beans. Five genera were identified: *Trichodorus*, *Helicotylenchus*, *Meloidogyne*, *Rotylenchulus*, and *Pratylenchus*. *Meloidogyne* spp had the highest population density compared to *Trichodorus*. According to Li *et al.* (2020), we can determine that the diversity of soil nematodes at a high level of taxonomy (for example, order) can be a more suitable indicator compared to diversity at a low level of taxonomy (for example, genus and family). A study by Maina *et al.* (2022) showed a high abundance of plant-parasitic in pigeon pea cultivated as a monocrop, indicating that appropriate methods for nematode management should be implemented. A study conducted by Ye *et al.* (2019) showed that after testing for species diversity in a population of root-knot nematodes, five species of root-knot nematodes were identified, namely *Meloidogyne incognita*, *Meloidogyne hapla*, *Meloidogyne partityla*, *Meloidogyne marylandi*, and *Meloidogyne haplanaria*. *Meloidogyne incognita* was the most abundant species in 95% of the samples, particularly in field crops such as cotton and soybeans (Ye *et al.*, 2019).

A survey conducted by Basavaraj *et al.* (2023) in India, following incidences of root-knot nematodes in common beans, found that the major symptoms include yellowing, root galls, root and tip rot, and marginal leaf drying. According to research (Indarti *et al.*, 2023), three nematode PPN genera were identified: *Helicotylenchus* spp.,

Meloidogyne spp., and *Rotylenchulus spp.* *Rotylenchulus spp.* was the most common. The C/N ratio had a positive effect on the population of *Meloidogyne spp.*, while total-N had a negative impact on the same species. It showed that the habitat conditions were homogeneous. A survey conducted by Krif *et al.* (2020) in Morocco aimed to determine the diversity and occurrence of PPNs in organically farmed fields. Twelve genera of PPNs were identified in root and soil samples. The most prevalent nematodes were root-lesion nematodes and root-knot nematodes. The bean crop exhibited high diversity indices of nematodes, particularly in terms of the plant-parasitic index, Shannon-Wiener index, and Evenness (Krif *et al.*, 2020).

A study conducted by Nisa *et al.* (2021) assessed the effects of edaphic and ecological factors on the diversity of soil nematodes and their structures at five landscape patches from five different sites in India. Among the patches, forest soil had the highest diversity, while alpine soil had the lowest; however, bacterivorous organisms were observed to dominate in all soil patches (Nisa *et al.*, 2021). An abundance of nematodes decreased from basic to acidic soil pH. Soil nutrients, such as phosphorus and nitrogen, were detrimental to nematode richness, particularly at every site, where nematode diversity was high in soils with abundant P and N but decreased with reduced soil nutrients (Lu *et al.*, 2020). As noted by Xiao *et al.* (2021), nematode diversity and richness reach their peak in tropical ecosystems. Carnivores/Omnivores and bacterial feeders had the least relative abundance, but high diversity in the tropical ecosystem, with an opposite pattern evident for plant and fungal feeders. According to Atandi *et al.* (2022), it can be determined that organic farming systems play a vital role in population buildup and improving the biodiversity of free-living nematodes compared to other systems. Soil properties significantly impact plant-parasitic nematode communities, particularly in organic farming systems (Krif *et al.*, 2020).

2.2.2 The Effect of Farmyard Manure, Chemical Fertilizer, and *Trichoderma harzianum* on Soil Nematodes under Common Beans

According to Termorshuizen *et al.* (2011), organic amendments enhance soil biological traits and suppress plant-parasitic nematodes. The major organic matter types used include composts, green manures, crop residues, agro-industrial waste, urban organic waste, animal manures, plant extracts, oil cakes, processing residues,

and peats. Suppression involves either an indirect or a direct mechanism (Oka 2010). Direct mechanisms include reducing the population of phytonematodes by releasing nematostatic, nematicidal, or allelochemical compounds mostly from organic matter. Indirect mechanisms include increasing diversity and population of antagonistic microorganisms (Moosavi, 2022). When soil is organically amended, there is an increase in the population of phytonematodes when environmental conditions favor parasitic nematodes. Hence, exploring the potential function of organic amendments, such as cattle manure, controls parasitic nematodes (Maina *et al.*, 2020). In the view of Atandi *et al.* (2017), who examined the comparison of various farming systems, organic farming, conventional farming, and farmer practice systems, and the differences in the suppression of PPNs in Kenya's maize and bean cropping systems. The study concluded that organic farming paradigms, especially those that employed the use of compost alongside Tithonia, remarkably helped reduce the abundance of *Meloidogyne* spp. and *Pratylenchus* spp. nematodes in soil. The management of root-knot nematodes using a bionematicide, such as *Trichoderma harzianum*, organic matter, and a soil fumigant has shown effective suppression of nematodes by limiting the application of chemicals (d'Errico *et al.*, 2022).

A study by Puissant *et al.* (2021) conducted a global analysis of 103 publications to assess the impact of agricultural practices on nematodes, which are key indicators of soil health. The study quantified the impact of every agricultural practice change on nematode indices. Globally, nematicides and organo-mineral fertilization showed the highest effect sizes, while mineral fertilization had the least. As noted by Li *et al.* (2022), organic fertilization promotes fungal and bacterial feeders at the level of trophic groups. At the community level, nematicide application reduces the abundance of nematodes and alters the structure of the food web, thereby encouraging copiotrophic nematode communities. Control of plant-parasitic nematodes using a biological agent called *Trichoderma* has been widely studied. Yield losses and plant death resulted from the high population of plant-parasitic nematodes. According to Almeida *et al.* (2022), *Trichoderma harzianum* and *Trichoderma asperellum* were compared to suppress plant-parasitic nematodes. *Trichoderma harzianum* reduced the population of nematodes compared to *Trichoderma asperellum*, which did not even promote plant growth. 1, 3-dichloropropene, a fumigant, was combined with organic fertilizer and *Trichoderma harzianum* to investigate its effect on plant yield, growth,

and root-knot nematode (d'Errico *et al.*, 2022). Nematode infestation was reduced in all crops by all treatments, with the greatest nematicidal effect observed in the mixture of the three products. Fumigant integration with *Trichoderma harzianum* and organic fertilizer increased yield and crop growth.

Besides plant-parasitic nematodes, common beans are associated with free-living nematodes. In the view of Andrassy (2025), free-living nematodes form the highest population of nematodes, yet they are given little attention. Free-living nematodes occupy various trophic levels and are grouped according to their feeding habits. Some free-living nematodes include omnivores, bacterial feeders, predatory nematodes, and fungal feeders (Yeates *et al.*, 1993). The roles of FLN are biocontrol of nematodes, organic matter decomposition, and nutrient mineralization; thus, they indicate the health status of the ecosystem (Ferris *et al.*, 2012). Bacterial feeders consume bacteria and contribute 30-50% of nitrogen in plants (Ingham *et al.*, 1985).

Fungal feeders puncture the cell wall of fungal hyphae using their stylet. Predatory nematodes consume invertebrates such as rotifers, protozoa, and other nematodes. Omnivores feed on fungi, plants, and invertebrates. Free-living nematodes (FLN) may accelerate soil organic matter decomposition. Numbers of FLN increase rapidly when inorganic fertilizers are applied, especially fungal and bacterial feeders (Maina *et al.*, 2021). Plant-pathogenic nematodes have increased due to the increased application of potassium. Disease and pest control in amended soil is achieved by various mechanisms that operate via their effects on the pathogen, host plant, and soil. The mechanisms may include the accumulation of microbial metabolites, deleterious decomposition of end products, and increased tolerance and disease resistance (Hailu & Hailu, 2020).

2.2.3 Morphology of Nematodes

The morphology of nematodes is visible to us only after special procedures for extraction, as they are microscopic and require a microscope (Maggenti, 2020). These microscopes and procedures fully determine our facts of nematode morphology, because they limit the level at which we can study nematodes (Van *et al.*, 2017). These procedures can result in substantial errors due to incorrect artifact interpretation, which they may create (Fourie *et al.*, 2017). They are very small;

hence, the challenge is to identify and detect nematodes using a microscope (Eisenback & Hunt, 2009).

2.2.4 The body of the Nematodes

Soil nematodes have a body that is vermiform, tapering less or more strongly towards the posterior and anterior ends, and with no appendages capable of independent movement (Waggoner, 2020). The tail is the posterior end. The head is the anterior end, which is not offset from the body by a distinct neck; instead, it often blends smoothly into the body (Kiontke & Fitch, 2013). The anterior end is also referred to as the cephalic region, lip region, or simply the anterior end. The entire body is covered by a less or more impermeable, relatively tough and elastic cuticle (Fourie *et al.*, 2017). The openings on the cuticle are for reproduction, "excretion," and digestion, as well as those of several secretory and/or sensory organs (Lazarova *et al.*, 2021).

2.2.5 Hatching and Life Cycle of Nematodes

The life histories of many parasitic and free-living nematodes are quite similar in that they have four larval stages (Mkandawire *et al.*, 2021). When the surroundings and temperature are unfavourable for the host, females of nematodes may not produce or produce few eggs (Wharton, 1986). Under less favourable conditions, such as extreme temperatures, a single female produces 300-500 eggs. However, in an optimum temperature of about 27 °C, more than 2800 eggs are produced (Luc *et al.*, 1990). Eggs can be laid stuck together or singly in masses in a gelatinous matrix secreted by females (Lee, 2002). A new generation emerges in 25 days, but if conditions are less favorable, development may end completely, or the life cycle could be extended to 30-40 days. Nematodes' lifespans vary from a few days to many years. During the dormant stage, each egg typically has a thick outer covering that protects it during the inactive period (Kumar & Yadav, 2020). Plant-parasitic nematodes are elongated, fusiform, slender, circular in cross-section, and tapering towards both ends. In root-knot nematodes, eggs are laid in egg sacs that are mostly buried within the host-derived root galls, a characteristic of *Meloidogyne* spp. induce through feeding (Coulibaly *et al.*, 2025).

Cysts and egg sacs protect eggs from natural enemies and desiccation. The juvenile in the egg matures into an adult by four moults. The first moult occurs in the egg. The egg develops into the first juvenile stage (J1) (Luc *et al.*, 1990). Juvenile coils numerous times within the eggshell and lies still. The J1 grows through the egg's first moult, then hatches as J2 (Fourie *et al.*, 2017). J2 is small in size, lacks organs of reproduction, and is fully developed. J2 undergoes a second moult and develops into J3, then a third moult to become J4 (Hooper *et al.*, 2005). The J4 stage goes through the fourth moult, and differentiation occurs, developing into adult males and females, which then mature. Temperature is important for the life cycle of nematodes, as illustrated by Agrios (2005). A life cycle from egg to adult can be completed in 3 to 4 weeks under optimal environmental conditions, such as 27°C and a soil pH of 4.0–8.0. The life cycle of *Ditylenchus dipsaci* takes 19- 23 days, whereas *Longidorus* spp. takes two years (Fourie *et al.*, 2017). Second-stage juveniles penetrate plants at a temperature between 10°C and 35°C, with 27°C being the optimum temperature, depending on the species.

In various nematode species, the life cycle of the nematodes is closely synchronized with that of the host, facilitated by host- and environment-derived stimuli, to enhance nematode reproduction (Kumar & Yadav, 2020). Each egg has a single juvenile, which hatches through cutting the eggshell with its stylet, through striking it with rhythmic blows that are intermittent or egg-shell rupturing with the tip of the tail, like in *Heterodera iri*, or by normal egg-shell rupture because of secretions of juvenile enzymes and movement (Mkandawire *et al.*, 2021). Eggs of cyst nematodes either survive in lemon-shaped (*Heterodera*) or round (*Globodera*) cysts in the soil, each with hundreds of eggs. Cysts have tiny openings at the vulval ends and neck where hatched juveniles can escape (Fourie *et al.*, 2017).

Once they hatch, the J2s of *Globodera pallida* and *Globodera rostochiensis* have a short survival span of less than two weeks without feeding, in comparison to *Tylenchulus semipenetrans* and *Meloidogyne javanica*, which can survive in the field for several months (Perry & Clarke, 1981). The occurrence of plant-parasitic nematodes has been reported to show a close association with leguminous crops, including common beans, as highlighted by Kimenju *et al.* (1999). Several nematode groups are commonly linked to beans: lesion nematodes such as *Aphelenchus* spp.,

Pratylenchus spp., *Criconebella spp.*, and *Tylenchus spp.*; root-knot nematodes (*Meloidogyne spp.*); stubby root nematodes (*Trichodorus spp.*); sheath nematodes (*Hemicycliophora spp.*); and other nematodes are associated with beans. *Meloidogyne spp.* are the most important because they are widely distributed, primarily in warm areas of the world, and exhibit a polyphagous nature (Coulibaly *et al.*, 2025).

2.3 The Effects of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard

Manure on Common Bean Growth and Yield Components.

A study by Tadesse & Abera (2023) found that applying farmyard manure increased yield components and the yield of haricot bean. Parameters like plant height, flowering, maturity, seed weight, and number of branches were influenced by farmyard manure. Research by Singh *et al.* (2022) found that integrating bio fertilizer and farmyard manure with 60 kg ha^{-1} single super-phosphate improved the yield of black gram by 3.28% and 7.4% respectively, compared to using 60 kg ha^{-1} single super-phosphate alone. Uptake of phosphorus in black gram when biofertilizer and FYM were applied together with single superphosphate showed improvement of uptake compared to 60kg ha^{-1} single superphosphate. According to the research by Shadrack *et al.* (2021), a study was conducted where Diammonium phosphate (0 and 62.5 kg ha^{-1}), Mavuno (0 and 185 kg ha^{-1}), lime (0 and 1.6 or 2.0 tha^{-1}), and Sympal (0 and 125 kg ha^{-1}) were applied under common beans. Applying lime and Mavuno increased common bean grain yields compared to the Sympal, control, and Diammonium phosphate. According to Raihan *et al.* (2021), farmyard manure contributed significantly to high seed germination rates, maximum bean height, accelerated first flowering days, increased number of pods per plant, maximum pod length of individual plants, pod weight, and pod width. Farmyard manure influenced the yield component and plant growth of the country bean.

A study by El-Yazal (2020) treated broad beans with different rates of minerals (NPK) and organic manure (farmyard). The plants that were organically fertilized attained higher values of various growth parameters (number of branches, plant height, dry weight, and plant leaflets' blade area) and yields, as well as broad bean components (number of pods/seeds, weight of pod/ seeds, number of plants/ pods, and weight of plant/seeds). According to the research (Nigatie, 2021), it can be determined that P and N fertilizers have a significant impact on common bean

production. Increasing nitrogen rates resulted in increased grain yields in two cultivars. The Dume variety showed great performance, followed by Nasir, while Ibbado indicated the least expectation. Research by Naluzze (2022) evaluated the effect of manure and NPK fertilizer on yield components of common beans. The inorganic fertilizer used was NPK 17:17:17; the results showed no significant effect on the growth parameters. Research conducted by Zamukulu *et al.* (2023) in eastern Democratic Republic of Congo determined that NPK fertilizer increased grain yield. It was evident in plants with high plant density compared to plants with low plant density. The study by Atandi *et al.* (2017) showed that organic farming paradigms, especially those that employed the use of compost alongside Tithonia, remarkably helped boost the growth of beans and elevate soil health.

Research conducted by Upenji (2020) over two seasons found that manure was applied at rates of 0 and 5 tons/ha. The seed yield of bean varieties significantly fluctuated according to manure input, variety, and season. The second season was more productive than the first season. The application of manure appeared to be the primary option for improving the seed yield of beans, thereby ensuring food security. According to Obsa (2017), it can be determined that the highest biomass and grain yield were achieved by applying the recommended P-fertilizer compared to vermicomposting material. Combining P-fertilizer and vermicomposting material increased yield compared to P-fertilizer alone, which can be considered an alternative for mostly integrated inorganic and organic nutrient sources for crop productivity and soil health. Plants fertilized with farmyard manure had higher values of Chlorophyll concentration, total amino acids, carotenoids, total carbohydrates, crude proteins, total sugars, insoluble carbohydrates, and reducing sugars in seeds and leaves compared to plants fertilized with NPK (El-Yazal, 2020).

A study by Ali *et al.* (2023) where NPK and organic fertilizers at a rate of 600 kg ha⁻¹ were applied to common beans. Organic fertilizers increased growth parameters (plant height and number of branches), yield components (number of seeds, grain yield, and number of pods), and dry matter compared to the control in common beans. According to Zongo *et al.* (2024), who determined the effects of applying organic compost and *Trichoderma harzianum* on the growth of soybeans and soil fertility. The analysis revealed that the application of organic compost mixed with *Trichoderma*

increased soil fertility by enhancing the presence of nitrogen (N), phosphorus (P), and potassium (K), which are vital in catalysing plant growth. Together, these practices demonstrated improved plant growth and nodulation, indicating that organic amendments and biocontrol agents can enhance nutrient availability and reduce the population of nematodes in the legume system (Zongo *et al.*, 2024).

The benefits of *Trichoderma* species include promoting growth, enhancing seed viability and germination, increasing flowering, photosynthesis, and yield, as well as improving root condition and structure (Tyskiewicz *et al.*, 2022). Mahmoodian *et al.* (2022) reported that different *Trichoderma harzianum* species have been used mostly as antagonists against various plant diseases. A direct relationship exists between fungi, biocontrol potential, and the rate at which enzymes degrade pathogens' cell walls. Chimeric chitinase 42 has been produced through protein engineering in various works by the author; the resultant protein was then transferred to *Trichoderma harzianum*. The research stability and presence of chimeric Chitinase were evident when the biocontrol potential and growth stimulation of six isolates were evaluated on common bean plants in greenhouse and laboratory conditions, compared to the wild type (Mahmoodian *et al.*, 2022). Seven *Trichoderma* species demonstrated increased shoot weight, root length, and shoot length in green gram (Mukhtar *et al.*, 2021).

Research by Elkelany *et al.* (2021) reported that three species of *Trichoderma*, *Trichoderma virens*, *Trichoderma viride*, and *Trichoderma harzianum*, improved the growth and yield of peanuts. The results were demonstrated by the dry and fresh weights of peanut plants, the weight of pods, and the number of pods. According to El-Nagdi *et al.* (2019), we can determine that after the application of *Trichoderma harzianum* in the soil, the growth and yield of pea also increased, which was demonstrated by the pod's dry weight, root fresh and dry weight, and shoot length. According to the research (Yao *et al.*, 2023), *Trichoderma* prevents diseases in plants, improves the utilisation of nutrients, and promotes growth. According to Olabiyi *et al.* (2016), we can determine that *Trichoderma harzianum* contributes to the effective growth of crops and a reduction in damage caused by nematodes.

2.4 Effect of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Plant-parasitic Nematodes in the Roots of Common Beans.

2.4.1 Nematode pathology

Plant-parasitic nematodes are biotrophic parasites; they obtain nutrients from the leaf cells, stem, and cytoplasm of living roots for growth, development, and survival. Nematodes have developed diverse feeding relationships and parasitic strategies with their plant hosts (Afzal & Mukhtar, 2024). They possess protrusible and hollow feeding structures, pharynxes, and stylets, and have undergone physiological and morphological adaptations to suit their feeding relationships (Eves-van den Akker, 2021). Depending on the species, they can feed on the cytoplasm of living plant cells that are either unmodified or have undergone modification to develop into elaborate feeding cells, as seen in root knot nematodes (Perry & Moens, 2011). The stylet of the nematode is used to penetrate and pierce the cell wall of the plant cell. Gland secretions are injected via the stylet orifice into the cell, where they are withdrawn as well as ingested by the nematode from the cytoplasm.

The stylet is also used by nematodes that enter the root tissue through openings or inject secretions to weaken or dissolve the middle lamella or cell wall (Arley Rey Páez, 2023). In general, all plant-parasitic nematodes damage plants through direct mechanical injury by use of the stylet in secretion and/or penetration of enzymes in the cells when the parasitic nematode is feeding (Bird & Bird, 2012). The physical presence of endoparasitic nematodes within the host typically disrupts the normal functioning of that host. However, the nematode architecture, feeding, and root system extent have changed, as it takes up water and nutrients from the soil less efficiently (Lee, 2002). The damage caused by nematodes depends on the inoculum density, that is, the infestation level. Moderate or low nematode numbers may result in more injury, but extremely high numbers can severely kill or damage their host.

2.4.2 Classification of Root-Knot Nematodes, *Meloidogyne* spp.

Root-knot nematodes belong to the kingdom; *Animalia*, phylum; *Nematoda*, class; *Nemata*, subclass; *Sercenentea*, order; *Tylnchida*, suborder; *Tylenchina*, family; *Meloidogynidae*, and genus; *Meloidogyne* (Perry *et al.*, 2009). There are 51 *Meloidogyne* species, of which *M. hapla*, *M. arenaria*, *M. javanica*, and *M. incognita* are economically significant in the production of beans worldwide (Churamani,

2017). Populations of root-knot nematodes consist of females and males, which are distinguished morphologically. Males are like worms and are 1.20- 1.50 mm long and have a diameter of 30- 60 μ m (body width) (Meza *et al.*, 2016). Female adults are pear-shaped and are 0.40 - 1.30 mm long by 0.27 - 0.75 mm in width. Second-stage juveniles have a vermiform shape, whereas third as well as fourth-stage juveniles are microscopic in size and sausage-shaped (Perry *et al.*, 2009).

2.4.3 Effect of Plant-parasitic Nematodes on Nodulation of Bean Roots

Plant-parasitic nematodes disrupt the process of nodulation in common beans by directly damaging the root tissues and indirectly affecting the symbiotic relationship with rhizobia (Kimenju *et al.*, 1999). Plant-parasitic Nematodes create a feeding site inside the nodule in the vascular bundle and encourage the development of giant cells that disrupt the function of the nodules. This results in nodules having premature senescence, hence a reduced ability to fix nitrogen (Kimenju *et al.*, 1999). They also affect nodulation by competing for nutrients and ecological niches, as well as suppressing the formation of lateral roots, thereby reducing the number of nodule formation sites. This results in the early degradation of nodules due to infection (Faridah & Van der Maesen, 1997). According to the research by Zongo *et al.* (2024), the application of *Trichoderma harzianum* resulted in improved nodulation in soybeans.

CHAPTER THREE

MATERIALS AND METHODS

3.1 Location of Study

The study was conducted in Karingani Ward, Chuka Sub-County, Tharaka Nithi County. The experiment was carried out from April 2024 to October 2024. The study site was located at a latitude of 00° 18'37" South and a longitude of 037° 39'38.0" East, approximately 1324 m above sea level (asl). The mean temperature is about 20 °C. The county has a bimodal rainfall pattern, with the long rains falling from April to June and the short rains from October to December. On average, the area receives about 1200 mm of rainfall annually. The soils in this area are Nitisols, characterized as deep, well-drained, strongly weathered, with moderate to high fertility, and possessing good water-holding capacity (Jaetold *et al.*, 2006). The climatic conditions are favourable for common bean farming. The economic activities in this area are agriculture, commerce, and trade. Agriculture is primarily an economic activity in this area, where various crops are supported by the region's agro-ecological zones, including cash crops (such as coffee and tea) and food crops (beans, maize, millet, and green grams). Dairy farming is also widely practiced, contributing significantly to household incomes.

3.2 Research Design and Treatments

The experimental design was a randomised complete block design (RCBD) with four treatments and three replications. The four treatments were: NPK (23.23.0) at 125 kg/ha, farmyard manure at 10 ton/ha as recommended by KARLO, *Trichoderma harzianum* (45g/20 l) as recommended by the manufacturer, and a control was assigned to three blocks. The plot size was 1.8 m by 1.5 m with a spacing of 0.45 m by 0.15 m, translating to 148,149 plants per hectare and 40 plants per plot. The spacing between plots was 1m. The experiment was conducted in two trials, each with a similar layout.

The layout was as follows:

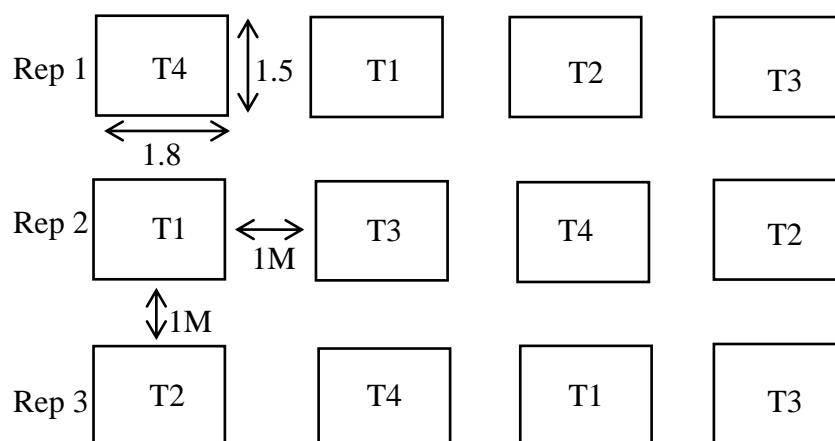


Figure 1: Farm Layout

Legend: T1-Control; T2-NPK (23.23.0) at 125 kg/ha; T3-farmyard manure at 10 tons/ha; T4-*Trichoderma harzianum*; Rep- Replicate

3.3 Crop Establishment and Management

Land preparation; the land was prepared by clearing previous crop residues and removing stumps and other foreign materials. Ploughing was done to a fine tilth, soil clods were broken, and land planning was considered, taking into account the type of irrigation used. The experimental field layout was done as per the above specifications. Planting; common bean seeds were sown in rows at a spacing of 0.45 x 0.15 m at two seeds per hill to a depth of 1cm, and requisite treatments applied per plot at sowing. Certified common beans were obtained from KALRO. Weeding; regular control of weeds was maintained to prevent competition for nutrients, water, space, and sunlight. Weed control was done manually by uprooting and using a panga. The appropriate method was uprooting because there was no damage to the roots. Weeds can harbour pests and diseases; therefore, weeding was done appropriately. Common pests and diseases were managed appropriately. Confidor WG 70 was drenched into the soil at a rate of 5g per 10 litres, as recommended by the manufacturer, to prevent cutworms from cutting germinating seedlings. Thinning; thinning was done two weeks after seedling emergence, at the four-leaf stage, when dried and weak seedlings were replaced. Irrigation; irrigation was done within the experimental site using a watering can. This method is good due to its water use

efficiency and small area of operation. The amount of water applied per plot was 40 litres. The irrigation was done in the evening as needed.

3.4 Sampling Procedures

3.4.1 Soil Sampling Procedure

The soil samples were taken from the field prior to sowing, at the flowering stage, and after harvesting the common beans in each trial. The soil sampling pattern was zig-zag to a depth of 25 cm with a soil auger at 4 points per plot. A composite sample of 300g was taken from each plot, packed in a zip bag, and stored in a cool box for nematode analysis in the lab.

3.4.2 Laboratory Procedure for Nematode Isolation (Modified Baermann Technique)

As described in Van (2006) nematode extraction, counting, and identification were carried out at the ICIPE Nematology laboratory. The Baermann technique was used to extract nematodes from the soil and roots. The procedure of nematode extraction from the soil was done as follows. 100 ml of soil was placed on the cotton-wool milk filter, which was held by support; maximum soil layer thickness was 2-3 mm. Support was placed with a sample into the funnel. Water was added from the side until the sieve's bottom touched the water. Nematodes left the soil, passed through the cotton-wool milk filter, and sank to the funnel's dish. Nematodes were collected after 48hours in Falcon tubes of 50 mL. Nematodes were allowed to settle on the Falcon tubes of 50 mL; supernatant was removed, passed through 25-micrometre sieves to reduce water volume. Nematodes were counted using a dissecting microscope at a magnification of 25-40x and identified using a compound microscope at a magnification of 10- 100x within a counting slide with a 2ml sample. Nematodes were identified to the genus level using an identification key and descriptions by Mai (2018)

3.4.3 Extraction of Nematodes from Common Bean Roots (Modified Baermann Technique)

Nematode extraction was done as described in Van (2006). Materials: knife, pair of scissors, cotton-wool milk filter, glass funnel with a piece of soft polyethene tube attached to the stem and closed with a screw clip, stand, support, 20 or 25µm sieve, 100ml glass beaker, dissecting microscope, and counting slide. Roots were chopped

into ± 1 cm pieces. 5g of roots were weighed using a weighing balance. Roots were placed on a cotton-wool milk filter placed within a support (sieve). Support with the sample was submerged gently in the water in the funnel. Nematodes were collected after 48 hours by opening the screw clip on the funnel stem. Nematodes were allowed to settle in the falcon tubes, and the suspension was passed over a 25 μm sieve to reduce the volume of water. Nematodes were counted using a dissecting microscope at a magnification of 25-40x and identified using a compound microscope at a magnification of 10- 100x within a counting slide with a 2ml sample. Nematodes were identified to the genus level using an identification key and descriptions by Mai (2018).

3.5 Data Collection

Data collection was done on common bean growth and yield components: height, number of leaves, number of branches (from the four middle tagged plants), emergence, biomass, number of pods, number of seeds, nodules, and grain yields. In contrast, nematode numbers were counted, and species diversity and evenness were calculated per plot. Data were collected under the following objectives.

3.5.1 To Determine the Effects of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Numbers, Evenness, and Species Diversity of Soil Nematodes under Common Beans.

Nematode species were identified up to the genus level and counted after carrying out the Baermann Technique in the laboratory (as described in 3.4.2).

Shannon Diversity Index (Konopinski, 2020)

$$H' = - \sum [(p_i) \times \ln (p_i)]$$

H'- Shannon diversity index

p_i - proportion of individuals of a nematode species in a whole community

\sum - sum symbol

ln- usually the natural logarithm (base $e \approx 2.718$)

Simpson's Index of Diversity (Kim et al., 2017)

$$D = 1 - \left(\frac{\sum n(n-1)}{N(N-1)} \right)$$

n- the total number of organisms of a particular nematode species

N- the total number of individual nematode species (all species combined)

The value D ranges from 0 to 1

Species Evenness (Mulder *et al.*, 2004)

$$EH = \frac{H'}{\ln(s)}$$

H'- Shannon diversity index

S- Total number of nematode species

In- Natural log

The value EH ranges from 0 to 1

3.5.2 To Determine the Effect of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Common Beans' Growth and Yield Components.

Common beans emergence; this was calculated as a percentage as the number of emerged seeds divided by the number of planted seeds multiplied by 100. Height; a ruler was used to measure the plant height from the base of the stem to the top of the canopy (this was the tip of the apical bud). It was done every two weeks and recorded in cm. Number of leaves; unfolded leaves of common beans were counted after every two weeks after sowing. All leaves were counted from the bottom of the crop. Number of branches; branches of common beans were counted after every two weeks after sowing. Branches were counted from the bottom of the crop. Biomass; destructive harvesting was done after maturity of common beans, where the plants were weighed after removing seeds from the pods using a weighing balance and recorded in kg. The samples were placed in an oven at 60°C for 48 hours in the Chuka University Plant Sciences laboratory. Dry biomass was weighed using a weighing balance and recorded in kg. Yield and yield components assessment; ten plants were randomly selected from each plot and tagged at pod maturity. Pods were harvested and placed in zip bags. The harvested pods from the sampled plants were shelled, and seeds counted for each plant. The average numbers of seeds per plant/plot were obtained. The final grain yield was determined by weighing all the seeds from the sampled plants and converting the yield to kilograms per hectare.

3.5.3 To Determine the Effect of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Plant-Parasitic Nematodes in the Roots of Common Beans

Root samples were taken from all plots with different treatments, i.e, NPK (23.23.0), farmyard manure, *Trichoderma harzianum*, and control. Nematode species were identified up to the genus level and counted after carrying out the modified Baermann Technique (as described in 3.4.3). Nodules; three plants were randomly selected from each plot and carefully dug out at the 70% flowering stage (45 days) after emergence. The plants were separated into shoots and roots. The roots were dipped in a bucket of water to remove the soil. The roots with undisturbed nodules were labelled in zip bags and then taken to the laboratory. Nodules were manually removed from the roots, and their numbers were recorded for each plant.

3.6 Data Analysis

Collected data on bean growth and yields were subjected to analysis of variance (ANOVA) using R version 4.5.1, and significant means were separated by LSD at $\alpha=0.05$. Species diversity was determined using Shannon and Simpson's indices, while evenness was calculated using Pielou's index on different treatments. The statistical model used is as follows:

$$Y_{ij} = \mu + \tau_i + \beta_j + \epsilon_{ij}$$

Y_{ij} : observed response (e.g plant height, biomass, number of leaves, seeds, pods, and branches) of the i^{th} treatment in the j^{th} replicate (block).

μ : this is the overall mean of all observations

τ_i : Effect of the i^{th} treatment where $i=1,2,3,4$

τ_1 - effect of control

τ_2 - effect of NPK 23.23.0

τ_3 - Effect of farmyard manure

τ_4 - effect of *Trichoderma harzianum*

β_j : effect of j^{th} block where $j=1,2,3$

ϵ_{ij} : random error term, assumed to be independently and normally distributed with mean 0 and constant variance.

3.7 Ethical Consideration

The Chuka University Institutional Ethics Review Committee requirements were followed keenly, ensuring confidentiality and security of collected data. An ethical clearance letter was acquired from Chuka University (Appendix 78). A permit was obtained from the National Commission for Science, Technology and Innovation (NACOSTI) after the proposal had been reviewed and approved (Appendix 79). All the collected data were used solely for academic purposes without reference to individuals.

CHAPTER FOUR
RESULTS AND DISCUSSION

4.1 Effects of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Numbers, Evenness, and Species Diversity of Soil Nematodes under Common Beans

4.1.1 Free-living Nematodes Identified in Soil

Table 1 presents the results from trials one and two, which determine the effect of different treatments (farmyard manure, *Trichoderma harzianum*, NPK, and control) on FLN populations at various growth stages of common beans (initial, flowering, and final). The FLN identified included predators, bacterivores, and *Aphelenchus spp*, *Dorylaimus spp*, and *Aphelenchoides spp*, and the total nematodes present (100ml soil sample).

Table 1: Free-living Nematodes Identified (100ml of soil) in NPK, FYM, and *Trichoderma harzianum* under Common Beans

| Trial | Stage | TRT | Bacterivores | Predator | <i>Aphelenchus spp</i> | <i>Aphelenchoides spp</i> | <i>Dorylaimus spp</i> | total |
|-------|-----------|---------|--------------|----------|------------------------|---------------------------|-----------------------|-------|
| 1 | Initial | Control | 200 | 65 | 280 | 13 | 57 | 615 |
| | | NPK | 107 | 97 | 144 | 30 | 81 | 459 |
| | | FYM | 100 | 5 | 106 | 0 | 12 | 223 |
| | | TH | 84 | 50 | 80 | 0 | 5 | 219 |
| | Flowering | Control | 100 | 6 | 138 | 0 | 33 | 277 |
| | | NPK | 87 | 0 | 78 | 23 | 0 | 188 |
| | | FYM | 167 | 11 | 123 | 0 | 15 | 316 |
| | | TH | 92 | 68 | 145 | 0 | 22 | 327 |
| | Final | Control | 75 | 4 | 77 | 0 | 0 | 156 |
| | | NPK | 63 | 0 | 60 | 10 | 0 | 133 |
| | | FYM | 206 | 15 | 200 | 0 | 35 | 456 |
| | | TH | 100 | 80 | 196 | 5 | 25 | 406 |
| 2 | Initial | Control | 75 | 4 | 77 | 0 | 0 | 156 |
| | | NPK | 63 | 0 | 60 | 10 | 0 | 133 |
| | | FYM | 206 | 15 | 200 | 0 | 35 | 456 |
| | | TH | 100 | 80 | 196 | 5 | 25 | 406 |
| | Flowering | Control | 17 | 8 | 30 | 8 | 0 | 63 |
| | | NPK | 26 | 0 | 60 | 15 | 0 | 101 |
| | | FYM | 584 | 0 | 162 | 0 | 85 | 831 |
| | | TH | 82 | 45 | 227 | 0 | 58 | 412 |
| | Final | Control | 10 | 13 | 40 | 0 | 0 | 63 |
| | | NPK | 104 | 23 | 105 | 60 | 25 | 317 |
| | | FYM | 983 | 34 | 312 | 0 | 97 | 1426 |
| | | TH | 163 | 100 | 312 | 100 | 70 | 745 |

Legend: TRT- Treatment; TH- *Trichoderma harzianum*; FYM- Farmyard manure; Initial- nematodes identified before applying treatments; Flowering- nematodes identified at flowering stage of common beans; Final- nematodes identified after harvesting common beans. *Aphelenchus spp*- fungal feeder; *Aphelenchoides spp*- fungal feeder; *Dorylaimus spp*- omnivores (some are fungal feeders).

Table 2: Effect of NPK, FYM, and *Trichoderma harzianum* on Free-living Nematodes under Common Beans

| TRIAL | STAGE | TREATMENT | FLN IN SOIL |
|-------|-----------|-----------------------|-------------------|
| 1 | Initial | Control | 615 ^a |
| | | NPK | 459 ^{ab} |
| | | FYM | 223 ^b |
| | | TH | 219 ^b |
| | Flowering | Control | 277 ^{ab} |
| | | NPK | 188 ^b |
| | | FYM | 316 ^b |
| | | TH | 327 ^a |
| | Final | Control | 156 ^b |
| | | NPK | 133 ^b |
| | | FYM | 456 ^a |
| | | TH | 406 ^a |
| 2 | Initial | Control | 156 ^b |
| | | NPK | 133 ^b |
| | | FYM | 456 ^a |
| | | TH | 406 ^a |
| | Flowering | Control | 63 ^c |
| | | NPK | 101 ^{bc} |
| | | FYM | 831 ^a |
| | | TH | 431 ^b |
| | Final | Control | 63 ^c |
| | | NPK | 317 ^c |
| | | FYM | 1426 ^a |
| | | TH | 745 ^b |
| | | Means | 370.75 |
| | | CV (%) | 25.47 |
| | | LSD _(0.05) | 183.25 |

Legend: CV- Coefficient of Variance; TH- *Trichoderma harzianum*; FYM- Farmyard manure; Initial- nematodes identified before applying treatments; Flowering- nematodes identified at flowering stage of common beans; Final- nematodes identified after harvesting common beans. Means with the same letters within the column are not significantly different at ($P < 0.05$).

As shown in Table 2 for trial one, at the initial stage, the control had the highest population of free-living nematodes (615), which was significantly different ($p < 0.05$) (Appendix 2) from TH (219) and FYM (223). At this stage, no treatment had been applied. At the flowering stage, TH (327) was significantly different ($p < 0.05$) (Appendix 3) from NPK (188) and FYM (316). Control treatment (277) was not significantly different ($p > 0.05$) from FYM, TH, and NPK. At this stage, FLN had begun to show improved decomposition of organic matter, while TH exhibited microbial interactions, which provided conducive conditions for FLN to thrive. At the final stage, TH (406) and FYM (456) were significantly different ($p < 0.05$) (Appendix 4) from NPK (133) and control (156), but there was no significant difference between TH and FYM. The difference between control, NPK, and organic treatments indicates

the availability of organic matter from FYM and TH, which supported FLN. Farmyard manure had the highest population of FLN, followed by TH, while control and NPK reduced FLN in both trials. Farmyard manure increased free-living nematodes because it has decomposable and organic matter, which stimulates the growth of FLN. *Trichoderma harzianum* is a fungus that suppresses PPNs but also encourages microbial biomass and diversity. This provides food sources and a favourable environment for free-living nematodes. Control treatment reduced FLN because natural resources were depleted.

In trial two, at the initial stage, TH (406) and FYM (456) were significantly different ($p < 0.05$) (Appendix 5) from NPK (133) and control (156), but there was no significant difference ($p > 0.05$) between TH and FYM. At the flowering stage, there was a notable increase where FYM (831) was significantly different ($p < 0.05$) (Appendix 6) from TH (431), NPK (101), and also Control (63). NPK and control were statistically similar. The results showed that FYM had a superior ability compared to TH, especially during the peak growth stage for common beans in supporting FLN. At the final stage, there were pronounced differences where FYM (1426) was significantly different ($p < 0.05$) (Appendix 7) from TH (745), NPK (317), and also Control (63).

Tables 1 and 2 show significant variations in free-living nematode population (FLN) across different treatments and bean growth stages. In trial two, farmyard manure (FYM) treatment demonstrated the highest nematode population, especially in the final stage, 456 (initial), 831 (flowering), 1426 (final), compared to the control, which had 156 (initial), 63 (flowering), and 63 (final). Application of FYM significantly enhances soil fertility and FLN biodiversity in common beans. This shows that organic amendments, such as FYM, provide conditions that favour nematode proliferation due to increased nutrient availability and microbial activity. In both trials and all stages of growth, FYM resulted in the highest populations of bacterivores, predators, *Aphelenchus spp*, and *Dorylaimus spp* nematodes compared to the control treatment especially in trial two. For instance in trial 1 FYM increased bacterivores from 100 (initial) to 167 (flowering) to 206 (final), predator from 5 (initial) to 11 (flowering) to 15 (final), *Aphelenchus spp* from 106 (initial) to 123 (flowering) to 200 (final) and *Dorylaimus spp* from 12 (initial) to 15 (flowering) to 35 (final). In

comparison to control in trial one which reduced bacterivores from 200 (initial) to 100 (flowering) to 75 (final), predator from 65 (initial) to 6 (flowering) to 4 (final), *Aphelenchus* spp from 280 (initial) to 138 (flowering) to 77 (final), *Aphelechoides* spp from 13 (initial) to 0 (flowering) to 0 (final) and *Dorylaimus* spp from 57 (initial) to 33 (flowering) to 0 (final).

These findings are in line with those of Yeates *et al.* (1993), who reported that organic amendments enhance the abundance and diversity of soil nematodes. This is evident in the final and flowering stages, where nematode numbers are elevated in comparison to other treatments. For instance, in trial two at the final stage, farmyard manure recorded 983 bacterivores. A High population of Bacterivores was attributed to organic matter decomposition in FYM, which is consistent with the findings of Ferris *et al.* (2001), who reported that organic amendments promote bacterivores nematodes. These findings compare to those of Atandi *et al.* (2022), who found that bacterial feeder nematodes were dominant FLN in systems that were organically managed compared to control and conventional systems. The presence of predator nematodes indicated a balanced soil ecosystem.

Predator nematodes contribute to the suppression of plant-parasitic nematodes and other soil organisms that can affect the yield of common beans. These findings corroborate with those of Kimenju *et al.* (2009), who reported that application of organic amendments suppresses PPNs by promoting beneficial microbial interactions. *Aphelenchus spp* are fungal feeders. When FYM is added to the soil, it results in increased fungal biomass since there is decomposition of organic matter. These fungal feeder nematodes play a key role in nutrient cycling and are an indicator of fungal activity in the rhizosphere of common beans. *Dorylaimus spp* have diverse feeding habits and act as omnivores. FYM resulted in a diverse microbial community, which provided a suitable environment for *Dorylaimus spp*. These findings comply with those of Hu *et al.* (2018), who reported that a high population of *Dorylaimus spp*, *Aphelenchus spp*, and predator nematodes was due to soil health and food resources in organically amended soil.

Trichoderma harzianum increased free-living nematodes in both trials, but trial two had the highest population. *Trichoderma harzianum* treatment increased progressively the populations of free-living nematodes from the initial to final stage, that is, bacterivores, predators, *Aphelenchus spp*, *Aphelenchoides spp*, and *Dorylaimus spp*, indicating a suppressive effect of biocontrol agents on plant-parasitic nematodes compared to control. For instance in trial two *Trichoderma harzianum* increased bacterivores from 100 (initial) to 82 (flowering) to 163 (final), predator from 80 (initial) to 45 (flowering) to 100 (final), *Aphelenchus spp* from 196 (initial) to 227 (flowering) to 312 (final), *Aphelenchoides spp* from 5 (initial) to 0 (flowering) to 100 (final) and *Dorylaimus spp* from 25 (initial) to 58 (flowering) to 70 (final) while control reduced bacterivores from 75 (initial) to 17 (flowering) to 10 (final), *Aphelenchus spp* from 77 (initial) to 30 (flowering) to 40 (final), *Aphelenchoides spp* from 0 (initial) to 8 (flowering) to 0 (final) and *Dorylaimus spp* from 0 (initial) to 25 (flowering) to 0 (final).

These results showed the role of TH in organic matter decomposition and soil microbial activity. *Trichoderma harzianum* is a fungus that colonizes soil organic residues, secretes extracellular enzymes, and accelerates the breakdown of organic substrates. This process releases nutrients and stimulates the proliferation of soil bacteria and fungi, which in turn provide abundant food resources for bacterivorous and fungivorous nematodes. Furthermore, *T. harzianum* contributes to improved soil biological quality by increasing microbial biomass, promoting nutrient mineralization, and enhancing soil enzyme activity. These processes enrich the soil food web and create favorable microhabitats that support a higher abundance of FLN. These findings corroborate with those of Sharon *et al.* (2011), who reported that *Trichoderma spp.* can suppress plant-parasitic nematodes (PPNs) through competition for resources and enzymatic degradation. The nematode population observed was consistent with these findings, showing its potential role in the management of parasitic nematodes.

These findings are consistent with those of Giada d'Errico *et al.* (2022), who investigated the combined application of *T. harzianum*, the fumigant 1,3-dichloropropene, and organic fertilizer on tomato crops infested with *Meloidogyne incognita*. The integrated treatment not only significantly reduced nematode

infestation but also enhanced plant growth and yield, suggesting that *T. harzianum* contributes to nematode suppression and promotes beneficial soil microbial communities. There was an increase in bacterivores, which are bacterial feeder nematodes. This increase resulted from *Trichoderma harzianum*, which enhanced soil microbial communities, hence promoting the abundance of bacterial feeder nematodes. *Aphelenchus spp* and *Aphelenchoides spp* increased because of an increase in fungal biomass in the soil after the application of *Trichoderma harzianum*. *Trichoderma harzianum* is also a fungus, and it has been shown to have altered the fungal community in favour of *Aphelenchoides spp.* and *Aphelenchus spp.* The findings are in line with those of Pinto *et al.* (2024), who reported that *Trichoderma spp.* not only protects plant roots from PPNs, but also influences fungal nematodes, which can make *Trichoderma* less effective in managing plant-parasitic nematodes. The effect of *Trichoderma harzianum* on *Dorylaimus spp.* varies. It showed a slight increase compared to the control treatment.

In comparison, control and NPK treatments had lower FLN populations compared to *Trichoderma harzianum* and FYM across all stages in both trials. For instance, in trial one NPK treatment resulted to reduced population of predator from 97 (initial) to 0 (flowering) to 0 (final), fungivorous; *Aphelenchus spp* from 144 (initial) to 78 (flowering) to 60 (final) and *Aphelenchoides spp* from 30 (initial) to 23 (flowering) to 10 (final) and omnivorous (*Dorylaimus spp*) from 81 (initial) to 0 (flowering) to 0 (final) and bacterivorous nematodes from 107 (initial) to 87 (flowering) to 63 (final). In comparison to control in trial one which reduced bacterivores from 200 (initial) to 100 (flowering) to 75 (final), predator from 65 (initial) to 6 (flowering) to 4 (final), *Aphelenchus spp* from 280 (initial) to 138 (flowering) to 77 (final), *Aphelenchoides spp* from 13 (initial) to 0 (flowering) to 0 (final) and *Dorylaimus spp* from 57 (initial) to 33 (flowering) to 0 (final). The NPK treatment, a chemical fertilizer, pollutes the soil, hence creating an unfavourable environment for FLN. They also do not contribute to microbial activities and organic matter decomposition, hence resulting to a reduced number of free-living nematodes, especially in the final stage. Control treatment means that no amendment was added to the soil, hence, organic matter decomposition was reduced. The soil was also not fertile, hence this contributed to a reduced number of FLN. Biomass in the soil was also reduced and, resulted to no degradation in the soil, hence reducing the number of FLN.

These findings compare to those of Neher (2010), who stated that chemical fertilizers reduced the abundance of free-living nematodes and organic amendments increased bacterivorous. These findings align with those of Ferris *et al.* (2001), who reported that chemical fertilizers reduced nematode diversity by having a negative impact on soil microbial communities. The effect of NPK on bacterivorous was variable and lower compared to FYM. These findings are consistent with those of Qi *et al.* (2023), who reported that nitrogen fertilizer increased the abundance of bacterivore nematodes while phosphorus fertilizer reduced the abundance of bacterivore nematodes. For instance, in trial two at the final stage, NPK had a higher population of bacterivores (104) than the control (10). This shows that when NPK provided essential nutrients for the growth of the plant, it may not have also provided a food source for bacterial nematodes like FYM. These findings are consistent with those of Roul *et al.* (2017), who reported that NPK fertilizer created uncondusive soil conditions, which affected the survival of predator nematodes due to an unstable food web, compared to soil treated with FYM. Natural soil conditions, i.e, plant root interactions and soil biological activity, contributed to the population of free-living nematodes identified in the control treatment.

4.1.2 Plant-parasitic Nematodes Identified in Soil

Table 3 presents the results from trial one and trial two, which determine the effect of different treatments (farmyard manure, *Trichoderma harzianum*, NPK, and control) on PPNs populations at various growth stages of common beans (initial, flowering, and final). The PPNs identified include *Tylenchus* spp, *Helicotylenchus* spp, *Meloidogyne* spp, *Trichodorus* spp, *Rotylenchus* spp, *Pratylenchus* spp, *Hoplolaimus* spp, together with the total nematode count.

Table 3: PPNs identified (100ml of soil) in NPK (23.23.23), FYM, and *Trichoderma harzianum* under Common Beans

| Trial | Stage | TRT | <i>Tylenchus</i> spp | <i>Helicotylenchus</i> spp | Meloidogyne spp | <i>Trichodorus</i> spp | <i>Rotylenchulus</i> spp | <i>Pratylenchus</i> spp | <i>Hoplaimus</i> spp | total |
|-------|-----------|-----|----------------------|----------------------------|-----------------|------------------------|--------------------------|-------------------------|----------------------|-------|
| 1 | Initial | CNT | 21 | 80 | 0 | 0 | 7 | 20 | 0 | 128 |
| | | NPK | 18 | 50 | 0 | 0 | 0 | 65 | 0 | 133 |
| | | FYM | 121 | 592 | 0 | 0 | 14 | 200 | 0 | 927 |
| | | TH | 64 | 276 | 0 | 0 | 100 | 110 | 0 | 550 |
| | Flowering | CNT | 30 | 100 | 0 | 0 | 25 | 36 | 0 | 191 |
| | | NPK | 24 | 108 | 0 | 0 | 0 | 67 | 0 | 199 |
| | | FYM | 42 | 466 | 0 | 0 | 0 | 150 | 0 | 658 |
| | | TH | 26 | 241 | 0 | 0 | 80 | 80 | 0 | 427 |
| | Final | CNT | 40 | 180 | 42 | 45 | 50 | 100 | 20 | 477 |
| | | NPK | 38 | 175 | 6 | 25 | 39 | 85 | 0 | 368 |
| | | FYM | 33 | 158 | 0 | 0 | 35 | 70 | 0 | 296 |
| | | TH | 23 | 137 | 0 | 0 | 7 | 70 | 0 | 237 |
| 2 | Initial | CNT | 40 | 180 | 42 | 45 | 50 | 100 | 20 | 477 |
| | | NPK | 38 | 175 | 6 | 25 | 39 | 85 | 0 | 368 |
| | | FYM | 33 | 158 | 0 | 0 | 35 | 70 | 0 | 296 |
| | | TH | 23 | 137 | 0 | 0 | 7 | 70 | 0 | 237 |
| | Flowering | CNT | 67 | 197 | 0 | 55 | 51 | 100 | 50 | 520 |
| | | NPK | 54 | 205 | 0 | 0 | 91 | 30 | 0 | 380 |
| | | FYM | 26 | 68 | 0 | 0 | 0 | 21 | 0 | 115 |
| | | TH | 0 | 31 | 0 | 7 | 5 | 10 | 0 | 53 |
| | Final | CNT | 85 | 402 | 0 | 130 | 158 | 168 | 125 | 1068 |
| | | NPK | 60 | 300 | 0 | 181 | 95 | 88 | 0 | 724 |
| | | FYM | 10 | 30 | 0 | 2 | 1 | 5 | 0 | 48 |
| | | TH | 0 | 4 | 0 | 0 | 2 | 3 | 0 | 9 |

Legend: CNT- control; TRT- Treatment; TH.- *Trichoderma harzianum*; FYM- Farmyard manure; Initial- nematodes identified before applying treatments; Flowering- nematodes identified at flowering stage of common beans; Final- nematodes identified after harvesting common beans

Table 4: Effect of NPK, FYM, and *Trichoderma harzianum* on Plant-parasitic Nematodes under Common Beans

| TRIAL | STAGE | TREATMENT | PPN IN SOIL |
|-------|-----------|-----------|-------------------|
| 1 | Initial | Control | 128 ^c |
| | | NPK | 133 ^c |
| | | FYM | 927 ^a |
| | | TH | 550 ^b |
| | Flowering | Control | 191 ^c |
| | | NPK | 199 ^c |
| | | FYM | 659 ^a |
| | | TH | 427 ^b |
| | Final | Control | 477 ^a |
| | | NPK | 368 ^{ab} |
| | | FYM | 296 ^b |
| | | TH | 237 ^b |
| 2 | Initial | Control | 477 ^a |
| | | NPK | 368 ^{ab} |
| | | FYM | 296 ^b |
| | | TH | 237 ^b |
| | Flowering | Control | 520 ^a |
| | | NPK | 380 ^a |
| | | FYM | 115 ^b |
| | | TH | 53 ^b |
| | Final | Control | 1064 ^a |
| | | NPK | 727 ^b |
| | | FYM | 48 ^c |
| | | TH | 9 ^c |
| | | Mean | 369.85 |
| | | CV (%) | 22.15 |
| | | LSD(0.05) | 182 |

Legend: CV- Coefficient of Variance; TH- *Trichoderma harzianum*; FYM- Farmyard manure; Initial- nematodes identified before applying treatments; Flowering- nematodes identified at flowering stage of common beans; Final- nematodes identified after harvesting common beans, Means with the same letters within the column are not significantly different at ($P < 0.05$).

As shown in Table 4, in trial one, in the initial stage, FYM (927) was significantly different ($p < 0.05$) (Appendix 11) from other treatments. *Trichoderma harzianum* (550) was significantly different ($p < 0.05$) from control (117) and NPK (133), which were not significantly different. At this stage, treatment had not been applied. At the flowering stage, as common beans continued to grow, counts of PPNs under FYM (659) were significantly different ($p < 0.05$) (Appendix 12) from NPK (199), TH (427), and control (191). *Trichoderma harzianum* had fewer PPNs compared to FYM, which showed its ability to suppress PPNs. At the final stage, control (477) was not significantly different from NPK (368), but it showed a significant difference ($p <$

0.05) (Appendix 13) from TH (237) and FYM (296). In trial two, at the initial stage, a similar trend to that observed in trial one was noted (Appendix 14). At the flowering stage, NPK (380) and control (520) were not significantly different, but they were significantly different ($p < 0.05$) (Appendix 15) from TH (53) and FYM (115). At the final stage, suppressive effects were pronounced. Control (1064) was significantly different ($p < 0.05$) (Appendix 16) from NPK (727), TH (9), and FYM (48).

Control increased PPNs because there were no soil amendments added to the soil to manage PPNs. The NPK treatment provided nutrients to common beans, hence promoted root growth and encouraged plenty hosts of PPNs. Chemical fertilizers also provide feeding sites for PPNs. The NPK treatment continued to increase PPNs in the soil by providing a conducive environment for PPNs. Use of NPK resulted in natural suppression of FLN by reducing organic matter decomposition and microbial activity. Chemical fertilizers like NPK also alters the soil conditions like pH, hence favouring PPNs compared to free-living nematodes. Farmyard manure contributed to organic matter decomposition, hence increasing FLN, which suppresses the PPNs. *Trichoderma harzianum* reduced the PPNs by promoting FLN and also acted as a biocontrol agent

As shown in Tables 3 and 4, there were distinct variations in PPNs across three stages in both trials. The highest PPN counts were observed in control and NPK treatments, especially in the final stage, while *Trichoderma harzianum* and FYM treatments significantly reduced PPNs population. Control treatment increased PPNs naturally because there were no amendments added. The NPK treatment encouraged plant growth, especially vigorous root growth, which provided feeding sites for PPNs, which increased the population of PPNs. Farmyard manure reduced plant-parasitic nematodes by encouraging free-living nematodes that suppress PPNs. *Trichoderma harzianum* suppressed PPNs through competition, parasitism, and producing nematicidal metabolites.

In trial two, control treatment showed a progressive increase in PPNs over time from 477 (initial) to 520 (flowering) to 1068 (final), NPK increased from 368 (initial) to 380 (flowering) to 724 (final), while *Trichoderma harzianum* reduced from 237 (initial) to 53 (flowering) to 9 (final). Farmyard manure treatment reduced PPNs from

296 (initial) to 115 (flowering) to 48 (final). Control increased populations of *Pratylenchus spp* from 100 (initial) to 100 (flowering) to 168 (final) and *Helicotylenchus spp* from 180 (initial) to 197 (flowering) to 402 (final), which were the dominant PPNs. Across both trials, in control, *Pratylenchus spp* increased from 20 (initial in trial one) to 168 (final in trial two). *Pratylenchus spp* are also known as root lesion nematodes, and their impact on common beans was root damage, causing lesions as well as reduced root growth. *Pratylenchus spp* reduced nutrient uptake since the roots were damaged. *Meloidogyne spp* was scarce and only found in the control, not at all sampling times, and only once at the final stage (42) in trial one. This is because they mostly live in roots, not soil, where eggs and females are inside the root, and you only catch free J2s, often few or none. Emergence of J2s are linked to temperature.

These findings compare to those of Abd-Elgawad (2020), who reported that soils with no amendment support increased populations of PPNs since there is no competition from nematode antagonists and beneficial microbes. Control had increased populations of *Tylenchus spp* from 40 (initial) to 67 (flowering) to 85 (final), *Meloidogyne spp* from 42 (initial) to 0 (flowering) to 0 (final), *Trichodorus spp* from 45 (initial) to 55 (flowering) to 130 (final), *Rotylenchulus spp* from 50 (initial) to 51 (flowering) to 158 (final), *Hoplolaimus spp* from 20 (initial) to 50 (flowering) to 125 (final) in trial two. In comparison to *Trichoderma harzianum*, which reduced *Tylenchus spp* from 23 (initial) to 0 (flowering) to 0 (final), *Helicotylenchus spp* from 137 (initial) to 31 (flowering) to 4 (final), *Trichodorus spp* from 0 (initial) to 7 (flowering) to 0 (final), *Rotylenchulus spp* from 7 (initial) to 5 (flowering) to 2 (final), and *Pratylenchus spp* from 70 (initial) to 10 (flowering) to 3 (final).

These findings corroborate with those of Ferris *et al.* (2001), who reported that a decline in organic amendments increases predation and microbial competition. *Hoplolaimus spp* were present in the control but absent in NPK treatments, because they are highly sensitive to chemical inputs and soil disturbances, aligning with findings by Neher *et al.* (2010), which showed that in trial two, *Hoplolaimus spp* populations increased from 20 (initial) to 50 (flowering stage) to 125 (final) in untreated soils. *Rotylenchulus spp* species were more prevalent in control plots, particularly at the final stage, reaching 158 PPNs. The NPK treatment had the second-

highest PPN population in both trials, especially at the final stage, where *Helicotylenchus spp* from 175 (initial) to 205 (flowering) to 300 (final), *Pratylenchus spp* from 85 (initial) to 30 (flowering) to 88 (final), and *Trichodorus spp* from 25 (initial) to 0 (flowering) to 181 (final) dominated compared to the control. These findings are consistent with those of Deepali (2023), who reported that synthetic fertilizers cause the PPNs population to increase by enhancing root growth, hence providing additional feeding sites. These findings are consistent with those of Ferris *et al.* (2001), who proposed that chemical fertilizers have a negative impact on nematode diversity by suppressing free-living nematodes like predators and *Dorylaimus spp*, indirectly favouring PPNs.

Farmyard manure substantially decreased PPN populations at all growth stages, leading to significantly lower final counts compared to the control treatment in both trials. For instance, in trial one, there was dominance of *Helicotylenchus spp* in the initial stage (592), which diminished drastically to 158 at the final stage. *Tylenchus spp* from 121(initial) to 42 (flowering) to 33 (final), *Rotylenchulus spp* from 200 (initial) to 150 (flowering) to 70 (final), also reduced drastically. These findings support research by Kimenju *et al.* (2009), which demonstrated that organic amendments like FYM improve soil microbial activity, enhancing antagonistic fungi and bacteria that suppress nematodes. Moreover, among all treatments, *Trichoderma harzianum* exhibited the most significant reduction in PPN populations compared to control treatment in both trials, with species such as *Tylenchus* from 23 (initial) to 0 (flowering) to 0 (final) and *Pratylenchus spp* from 70 (initial) to 10 (flowering) to 3 (final) nearly eliminated by the final stage in trial two. For example, in trial two, the initial *Helicotylenchus spp* population 137 (initial) declined sharply to just 4 (final) at the final sampling. *Trichoderma harzianum* showed a suppressive effect on plant-parasitic nematodes.

These results comply with those of Sharon *et al.* (2011), who highlighted that the biocontrol properties of *Trichoderma harzianum*, attributing its effectiveness to enzyme production that disrupts nematode eggs and juveniles. Furthermore, these findings are in line with those of Neher *et al.* (2010), who emphasized that biocontrol agents contribute to enhanced soil suppressiveness, limiting plant-parasitic nematode proliferation. In contrast to these findings, Lawal & Atungwu (2017) reported that

NPK fertilizer increased plant-parasitic nematodes compared to the control treatment, while organic fertilizer suppressed PPNs.

4.1.3 Effect of NPK (23.23.0), Farmyard manure, and *Trichoderma harzianum* on Shannon Diversity, Simpson's Index of Diversity, and Evenness of Nematodes

Table 5 presents results from two trials determining the effects of different treatments on the diversity and evenness of nematodes at two growth stages: final and initial. The treatments are control, NPK, farmyard manure, and *Trichoderma harzianum*.

Table 5: Effect of NPK (23.23.0), Farmyard manure, and *Trichoderma harzianum* on Shannon Diversity, Simpson's Index of Diversity, and Evenness of Nematodes

| Tri al | Stage | TRT | Free-living nematodes | | | Plant-parasitic nematodes | | |
|-----------|---------|-----|-----------------------|------------------------|--------------|---------------------------|------------------------|--------------|
| | | | Shannon diversity | Simpson's diversity | Even ness | Shannon diversity | Simpson's diversity | Even ness |
| 1 | Initial | CNT | 1.26 | 0.67 | 0.78 | 1.04 | 0.56 | 0.75 |
| | | NPK | 1.52 | 0.77 | 0.94 | 0.99 | 0.61 | 0.90 |
| | | FYM | 0.96 | 0.57 | 0.69 | 0.95 | 0.53 | 0.68 |
| | | TH | 1.16 | 0.67 | 0.84 | 1.23 | 0.66 | 0.89 |
| | Final | CNT | 0.79 | 0.53 | 0.72 | 1.71 | 0.78 | 0.88 |
| | | NPK | 0.91 | 0.57 | 0.83 | 1.41 | 0.70 | 0.79 |
| | | FYM | 1.03 | 0.60 | 0.74 | 1.17 | 0.63 | 0.85 |
| | | TH | 1.24 | 0.67 | 0.77 | 1.00 | 0.25 | 0.73 |
| 2 | Initial | CNT | 0.79 | 0.53 | 0.72 | 1.71 | 0.78 | 0.88 |
| | | NPK | 0.91 | 0.57 | 0.83 | 1.41 | 0.70 | 0.79 |
| | | FYM | 1.03 | 0.60 | 0.74 | 1.17 | 0.63 | 0.85 |
| | | TH | 1.24 | 0.67 | 0.77 | 1.00 | 0.25 | 0.73 |
| | Final | CNT | 0.91 | 0.54 | 0.82 | 1.65 | 0.78 | 0.92 |
| | | NPK | 1.44 | 0.74 | 0.89 | 1.44 | 0.73 | 0.90 |
| | | FYM | 0.86 | 0.47 | 0.62 | 1.07 | 0.56 | 0.66 |
| | | TH | 1.46 | 0.73 | 0.91 | 1.06 | 0.72 | 0.97 |

Legend: CNT- control; TRT- Treatment; TH- *Trichoderma harzianum*; FYM- Farmyard manure; Initial- before applying treatments and sowing; Final- harvesting common beans

The diversity of FLN and PPNs was calculated using Simpson's and Shannon indices, together with evenness. Ecological indices like Shannon Index, Simpson's index, and Evenness were used to calculate diversity as well as the distribution of PPNs and FLN. Shannon diversity focuses on evenness (how species are equally distributed) and species richness (the total number of species present). Simpson's Index of

diversity estimates the likelihood that two randomly chosen individuals from a given sample will be from different species. As presented in Table 5, different treatments had distinct effects on the diversity and evenness of nematodes.

As shown in Table 5, the Shannon diversity index of FLN showed an overall rise, especially *Trichoderma harzianum*, which had the highest increase in both trials. For example, in trial one, Shannon diversity of *Trichoderma harzianum* increased from 1.16 (initial) to 1.24 (final) compared to the control, which reduced from 1.26 (initial) to 0.79 (final). Farmyard manure didn't show a pronounced increase from 0.96 (initial) to 1.03 (final) in diversity because there was dominance of certain nematodes like bacterial feeders, as shown in Table 1. In trial one NPK reduced Shannon diversity of FLN from 1.52 (initial) to 0.92 (final). The NPK treatment created unfavourable conditions for FLN, as well as suppressing FLN. This imbalance reduced evenness, resulting in reduced diversity after application of NPK treatment. Simpson's index of diversity of FLN had a comparable trend. *Trichoderma harzianum* increased Simpson's index of diversity in trial two from 0.67 (initial) to 0.73 (final) compared to the control, which did not show a significant increase from 0.53 (initial) to 0.54 (final). These results show that there was a reduction in species dominance, resulting in a balanced nematode community because the control treatment did not increase the number of FLN, while *Trichoderma harzianum* increased. *Trichoderma harzianum* reduced dominant PPNs and suppressed them, resulting in higher evenness and a balanced community.

Evenness values of FLN remained relatively consistent across the different treatments, suggesting that species distribution did not change significantly despite increases in overall diversity. The *Trichoderma harzianum* treatment recorded the highest final evenness value (0.91) in trial two, reflecting a well-balanced distribution within the nematode community. In comparison, the control treatment had a final value of Evenness of 0.82. These results compare to those of Rigobelo *et al.* (2024), who studied the effects of *T. harzianum* on the diversity of root and soil microbiomes in soybean plants and found that soil samples treated with *T. harzianum* exhibited increased richness and diversity, as measured by both Shannon and Gini-Simpson indices, suggesting a positive impact on soil microbial diversity.

Shannon diversity index for plant-parasitic nematodes showed mixed responses across treatments. Control treatment showed the highest increase in the Shannon diversity index. For instance, in trial one, diversity increased from 1.04 (initial) to 1.71 (final). NPK showed a slight increase from 0.99 (initial) to 1.41 (final) while *Trichoderma harzianum* reduced from 1.23 (initial) to 1.00 (final). Control increased the diversity because no soil amendment was added, and no soil disturbance occurred. This made more PPNs coexist and survive without competition or suppression, resulting in high evenness. However, the *Trichoderma harzianum* from 1.23 (initial) to 1.00 (final) and FYM from 0.95 (initial) to 1.17 (final) treatments exhibited relatively stable values, suggesting that these treatments may not significantly impact plant-parasitic nematode diversity. Farmyard manure and *Trichoderma harzianum* reduced Shannon diversity of PPNs because they created conditions unfavourable for plant-parasitic nematodes. *Trichoderma harzianum* is a biocontrol, hence it suppressed the plant-parasitic nematodes while FYM promoted the FLN. This selection pressure resulted in reduced dominant species, hence a drop in diversity. The NPK treatment increased Shannon diversity from 0.99 (initial) to 1.41 (final) because it enhanced the growth of common beans, providing more nutrients and roots, which supported plant-parasitic nematodes. This also increased evenness, resulting in a high Shannon diversity.

In trial one, Simpson's Index of Diversity for plant-parasitic nematodes increased in the control from 0.56 (initial) to 0.78 (final) and NPK from 0.61 (initial) to 0.70 (final), implying reduced dominance by a few species. However, *Trichoderma harzianum* showed a decline in Simpson's index from 0.66 (initial) to 0.25 (final), suggesting that a few plant-parasitic nematode species may have become dominant under this treatment. Evenness values fluctuated across treatments, with *Trichoderma harzianum* showing a final increase from 0.73 (initial) to 0.96 (final) in trial two, suggesting a more uniform distribution of plant-parasitic nematodes. These findings are consistent with those of Dash *et al.* (2018), who reported that the use of NPK increased the nematode diversity and populations of PPNs. These findings are contrary to those of Qi *et al.* (2023), who reported that nitrogen and phosphorus fertilizers affected the diversity of soil nematode communities.

4.2 Effect of NPK (23.23.0), *Trichoderma Harzianum*, and Farmyard Manure on Common Beans' Growth and Yield Components.

4.2.1 Effect of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Common Beans' Seed Emergence

Seed emergence is a critical early stage of seed vigour and subsequent crop establishment in legumes, including common beans (*Phaseolus vulgaris*). Table 6 presents the percentage of seed emergence under four different treatments: NPK (23.23.0), farmyard manure, *Trichoderma harizianum*, and control, across two trials. In trial 1, *Trichoderma harzianum* had the highest emergence (92.5%), while in Trial 2, FYM led with 93.33%. Notably, all treatments were not significantly different ($p>0.05$) (Appendix 20 and 21) since they shared grouping letter "a" and the Least Significant Difference (LSD ($_{0.05}$)) value of 17.20, which exceeds the observed differences among treatments.

Table 6: Effect of NPK (23.23.0), Farmyard Manure, and *Trichoderma harizianum* on Seed Emergence of Common Beans.

| TRIAL | TREATMENT | Seed Emergence (%) |
|----------|------------------------------|--------------------|
| 1 | Control | 83.33 ^a |
| | NPK | 79.17 ^a |
| | Farmyard manure | 86.67 ^a |
| | <i>Trichoderma harzianum</i> | 92.50 ^a |
| 2 | Control | 84.17 ^a |
| | NPK | 84.17 ^a |
| | Farmyard manure | 93.33 ^a |
| | <i>Trichoderma harzianum</i> | 77.50 ^a |
| | Mean | 85.10 |
| | CV (%) | 16.59 |
| | LSD _(0.05) | 17.20 |

Legend: CV- Coefficient of variation, Means with the same letters within the column are not significantly different at ($P < 0.05$).

As presented in Table 6, in trial one, *Trichoderma harzianum* had the highest seed emergence of 92.50% compared to the control, which had 83.33%. In trial two, farmyard manure had the highest seed emergence of 93.33% compared to the control, which had 84.17%. Inconsistent seed emergence of *Trichoderma harzianum* in both trials, 92.50% in trial one and 77.50% in trial two, compared to the control, which had 83.33% in trial one and 84.17% in trial two, could have been caused by factors like different environmental conditions (moisture content and soil temperature) between

trials, which influence the efficacy of microbial inoculants. FYM had relatively good seed emergence in both trials, 86.67% in trial one and 93.33% in trial two, compared to the control, which had 83.33% in trial one and 84.17% in trial two. This indicates that germination was primarily determined by seed vigour and uniform germination conditions rather than treatment effects. Emergence was dependent on existing environmental and soil conditions, which were adequate for initial germination.

Despite lack of significant differences, these findings are contrary to those of Harman *et al.* (2004), who reported that *Trichoderma harzianum* has been shown to enhance seed emergence through mechanisms like improved nutrient availability, phytohormone production, and biocontrol of pathogens. The findings are contrary to those of Singh *et al.* (2024), who reported that organic amendments enhance soil structure and microbial activity, creating favourable conditions for plant development beyond germination. These findings are contrary to those of Yang (2017), who reported that the application of balanced NPK fertilizer increased the percentage and germination ability of seeds.

4.2.2 Effect of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Common Beans' Height

Table 7 presents the results on the height (in cm) of common bean crops recorded at 14, 28, 42, and 56 days after sowing (DAS) under different treatments across the two trials. The treatments included NPK fertilizer, control, *Trichoderma harzianum*, and farmyard manure. Each treatment showed a significant difference ($p < 0.05$) (Appendix 23 to 30) in height of common beans over time, where NPK had the tallest plants in both trials, followed by farmyard manure, *Trichoderma harzianum*, and control. CV and LSD ($\alpha = 0.05$) were used to show the significance and reliability of observed differences.

As shown in Table 7, there were significant differences ($P < 0.05$) (Appendix 23 to 30) among all treatments. Trial two had higher growth responses compared to trial one, but similar trends were observed in trial two. This showed that there was residual fertility, and soil conditions improved. Looking at the trends in both trials on the increase of plant height from 14 to 28 to 42 to 56 DAS, NPK had the highest increase, followed by FYM, then *Trichoderma harzianum*, and Control was the last. The NPK

treatment had the tallest plants in both trials compared to the control treatment and the other treatments. The NPK treatment supplied readily available nutrients, which accelerated growth, hence increased the height of common beans, while FYM slowly released nutrients for common bean uptake, improved water retention, soil structure, and microbial activity for plant growth. *Trichoderma harzianum* enhanced nutrient uptake but to a lesser extent compared to NPK and FYM, root health indirectly supporting common bean growth. In comparison to control, which depended solely on existing soil fertility, which limited common bean development and reduced overall growth performance.

Table 7: Effects of NPK (23.23.0), Farmyard manure, and *Trichoderma harzianum* on the Height of Common Beans.

| TRIAL | TREATMENT | HEIGHT (in cm) | | | |
|-------|------------------------------|--------------------|--------------------|--------------------|---------------------|
| | | 14 DAS | 28 DAS | 42 DAS | 56 DAS |
| I | Control | 8.25 ^d | 14.58 ^d | 18.92 ^d | 26.17 ^d |
| | NPK | 19.75 ^a | 35.17 ^a | 50.42 ^a | 66.67 ^a |
| | Farmyard manure | 15.75 ^b | 29.83 ^b | 39.17 ^b | 55.33 ^b |
| | <i>Trichoderma harzianum</i> | 12.75 ^c | 21.58 ^c | 29.83 ^c | 40.42 ^c |
| II | Control | 8.08 ^d | 10.58 ^d | 35.25 ^d | 52.00 ^d |
| | NPK | 16.25 ^a | 59.50 ^a | 96.67 ^a | 104.92 ^a |
| | Farmyard manure | 13.25 ^b | 30.25 ^b | 77.00 ^b | 85.67 ^b |
| | <i>Trichoderma harzianum</i> | 10.58 ^c | 19.58 ^c | 58.58 ^c | 67.42 ^c |
| | Mean | 13.07 | 27.64 | 50.72 | 62.32 |
| | CV (%) | 14.96 | 33.19 | 27.17 | 20.40 |
| | LSD _(0.05) | 1.123 | 5.263 | 7.908 | 7.296 |

Legend: CV- Coefficient of variation; DAS- Days after sowing. Means with the same letters within the column are not significantly different at (P <0.05).

As shown in Table 7, for instance, in trial two, NPK treatment increased height from 16.25 at 14 DAS to 59.50 at 28 DAS to 96.67 at 42 DAS to 104.92 at 56 DAS compared to control, which increased from 8.08 at 14 DAS to 10.58 at 28 DAS to 35.25 at 42 DAS to 52.00 at 56 DAS. The NPK treatment provided readily available macro-nutrients, that is, nitrogen and phosphorus, which are essential for the growth of common beans. Nitrogen is one of the key elements in shoot and leaf development. Phosphorus plays vital roles like helping in cell division, energy transfer, and root development. Farmyard manure was ranked second in both trials. For instance, in trial one, FYM increased plant height from 15.75 at 14 DAS to 29.83 at 28 DAS to 39.17 at 42 DAS to 55.33 at 56 DAS compared to the control, which increased from 8.25 at 14 DAS to 14.58 at 28 DAS to 18.92 at 42 DAS to 26.17 at 56 DAS. Plants in plots

treated with FYM were taller compared to those with no treatment and *Trichoderma harzianum*. Farmyard manure led to an increase in plant height because of the presence of organic matter, which improved soil fertility and provided essential nutrients gradually.

Trichoderma harzianum showed less pronounced results on plant height compared to NPK and FYM. For instance, in trial two, *Trichoderma harzianum* increased plant height from 10.58 at 14 DAS to 19.58 at 28 DAS to 58.58 at 42 DAS to 67.42 at 56 DAS compared to the control, which increased from 8.08 at 14 DAS to 10.58 at 28 DAS to 35.25 at 42 DAS to 52.00 at 56 DAS. However, *Trichoderma harzianum* encouraged growth compared to control treatment. *Trichoderma harzianum* increased the growth of common beans by producing growth regulators. Control increased plant height from 8.25 at 14 DAS to 14.58 at 28 DAS to 18.92 at 42 DAS to 26.17 at 56 DAS in trial one and increased from 8.08 at 14 DAS to 10.58 at 28 DAS to 35.25 at 42 DAS to 52.00 at 56 DAS in trial two. Control treatment had the shortest plants compared to all other treatments, this is because there were no amendments added to the soil.

These findings compare to those of Halifu *et al.* (2019), who showed that when *Trichoderma spp.* is applied in the plant's rhizosphere, it promotes morphological traits like height, number of branches, leaves, biomass, and shoot-root length. These findings corroborate with those of Uddin *et al.* (2023), who reported that NPK fertilizer increased the plant height compared to farmyard manure of garden pea. These findings are in line with those of Khan *et al.* (2021), who suggested that organic manures not only increased growth but also nutrient uptake compared to the control treatment in chickpea. These findings are consistent with Ali *et al.* (2023), who reported that organic fertilizers applied at the rate of 600 kg ha⁻¹ increased the plant height compared to the control in common beans. These results are inconsistent with those of El-Yazal (2020), who reported that FYM increased plant height compared to NPK and control treatment in broad beans. These findings contrast those of Hossain & Akter (2020), who reported that *Trichoderma spp.* increased growth compared to farmyard manure and control treatment, which had a lower growth rate of brinjal.

4.2.3 Effects of NPK (23.23.0), Farmyard manure, and *Trichoderma harzianum* on the Number of Leaves and Branches of Common Beans

Tables 8 and 9 present the results of the number of branches and leaves of common beans, respectively. There was a significant difference ($p < 0.05$) in the number of branches across all treatments in both trials. There was no significant difference ($p > 0.05$) in trial one between FYM and *Trichoderma harzianum* on 56 DAS, but there was a significant difference ($p < 0.05$) (Appendix 35 to 42) in trial two across all treatments in the number of leaves.

Table 8: Effects of NPK (23.23.0), Farmyard manure, and *Trichoderma harzianum* on Number of Branches of Common Beans.

| TRIAL | TREATMENT | NUMBER OF BRANCHES | | | |
|-------|------------------------------|--------------------|-------------------|--------------------|--------------------|
| | | 14 DAS | 28 DAS | 42 DAS | 56 DAS |
| I | Control | 1.58 ^c | 2.67 ^d | 3.92 ^c | 4.50 ^d |
| | NPK | 2.50 ^a | 8.58 ^a | 13.08 ^a | 15.00 ^a |
| | Farmyard manure | 2.08 ^b | 5.91 ^b | 8.58 ^b | 9.83 ^b |
| | <i>Trichoderma harzianum</i> | 2.00 ^b | 4.25 ^c | 6.58 ^c | 7.92 ^c |
| II | Control | 0.50 ^c | 2.25 ^d | 5.42 ^d | 7.00 ^d |
| | NPK | 1.42 ^a | 8.42 ^a | 28.08 ^a | 32.08 ^a |
| | Farmyard manure | 1.00 ^b | 5.58 ^b | 15.67 ^b | 20.75 ^b |
| | <i>Trichoderma harzianum</i> | 0.92 ^b | 3.42 ^c | 9.33 ^c | 12.41 ^c |
| | Mean | 1.5 | 5.14 | 11.33 | 13.69 |
| | CV (%) | 25.56 | 20.45 | 34.19 | 33.46 |
| | LSD _(0.05) | 0.22 | 0.60 | 2.22 | 2.63 |

Legend: CV- Coefficient of variation; DAS- Days after sowing, Means with the same letters within the column are not significantly different at ($P < 0.05$).

Table 9: Effects of NPK (23.23.0), Farmyard manure, and *Trichoderma harzianum* on the Number of Leaves of Common Beans.

| TRIAL | TREATMENT | NUMBER OF LEAVES | | | |
|-------|------------------------------|-------------------|--------------------|--------------------|--------------------|
| | | 14 DAS | 28 DAS | 42 DAS | 56 DAS |
| I | Control | 6.75 ^c | 10.00 ^d | 13.75 ^d | 15.50 ^c |
| | NPK | 9.50 ^a | 27.75 ^a | 41.25 ^a | 47.00 ^a |
| | Farmyard manure | 8.25 ^b | 19.75 ^b | 27.75 ^b | 31.00 ^b |
| | <i>Trichoderma harzianum</i> | 8.00 ^b | 14.75 ^c | 21.75 ^c | 25.75 ^b |
| II | Control | 3.50 ^c | 8.75 ^d | 18.25 ^d | 23.00 ^d |
| | NPK | 6.25 ^a | 27.25 ^a | 86.25 ^a | 98.25 ^a |
| | Farmyard manure | 5.00 ^b | 18.75 ^b | 49.00 ^b | 64.25 ^b |
| | <i>Trichoderma harzianum</i> | 4.75 ^b | 12.25 ^c | 30.00 ^c | 39.25 ^c |
| | Mean | 6.5 | 17.41 | 36 | 43 |
| | CV (%) | 17.70 | 18.10 | 32.29 | 32.08 |
| | LSD _(0.05) | 0.66 | 1.81 | 6.67 | 7.92 |

Legend: CV- Coefficient of variation; DAS- Days after sowing, Means with the same letters within the column are not significantly different at (P <0.05).

The number of leaves and branches is a generally critical vegetative characteristic that contributes to the architecture, reproductive potential, and biomass production. The NPK treatment showed the highest number of leaves and branches in both trials, followed by farmyard manure, *Trichoderma harzianum*, and control. The trends observed in both trials were similar for the number of leaves and branches. Trial two had a higher number of leaves and branches compared to trial one this is because there was residual fertility, and soil conditions improved.

As shown in Tables 8 and 9, looking at the trends in both trials on the increase of the number of branches and leaves from 14 to 28 to 42 to 56 DAS, NPK had the highest increase, followed by FYM, then *Trichoderma harzianum*, and Control had the least. The NPK treatment supplied readily available nutrients, which accelerated growth, hence the highest number of leaves and branches, while FYM slowly released nutrients for common bean uptake, improved water retention, soil structure, and microbial activity for plant growth. *Trichoderma harzianum* enhanced nutrient uptake but to a lesser extent compared to NPK and FYM, root health indirectly supporting moderate growth. In comparison to the control, which depended solely on existing soil fertility, which limited common bean development and reduced overall growth performance.

For instance, in trial one, NPK fertilizer increased the number of branches of common beans from 2.50 at 14 DAS to 8.58 at 28 DAS to 13.08 at 42 DAS to 15.00 at 56

DAS, while control increased from 1.58 at 14 DAS to 2.67 at 28 DAS to 3.92 at 42 DAS to 4.50 at 56 DAS. Table 9 shows that NPK, for instance, in trial one, increased the number of leaves from 9.50 at 14 DAS to 27.75 at 28 DAS to 41.25 at 42 DAS to 47.00 at 56 DAS, while control increased from 6.75 at 14 DAS to 10.00 at 28 DAS to 13.75 at 42 DAS to 15.50 at 56 DAS. The NPK treatment provided essential macronutrients, that is, nitrogen and phosphorus, which play key roles in processes of vegetative growth like cell differentiation, elongation, and cell division. Nitrogen played a key role in shoot growth as well as new branches. Nitrogen made leaves greener because of increased chlorophyll that was produced. Phosphorus aids in root development and energy transfer, which promotes shoot proliferation. Phosphorus also improves the strength of branches and stems and overall health.

Farmyard manure also influenced the number of leaves and branches compared to *Trichoderma harzianum* and the control in both trials. For example, in trial two, FYM increased the number of branches (as shown in Table 6) from 1.00 at 14 DAS to 5.58 at 28 DAS to 15.67 at 42 DAS to 20.75 at 56 DAS, while the control increased from 0.50 at 14 DAS to 2.25 at 28 DAS to 5.42 at 42 DAS to 7.00 at 56 DAS. As presented in Table 7 in trial two, FYM increased the number of leaves from 5.00 at 14 DAS to 18.75 at 28 DAS to 49.00 at 42 DAS to 64.25 at 56 DAS compared to control 3.50 at 14 DAS to 8.75 at 28 DAS to 18.25 at 42 DAS to 23.00 at 56 DAS. Farmyard manure was less effective compared to NPK. These findings concur with earlier findings, which state that organic amendments like farmyard manure release nutrients gradually and are influenced by microbial mineralization for nutrients to be available for the plant. The increase in the number of leaves and branches over time showed that there was a gradual release of nutrients that sustained vegetative growth. This was evident on the last day of data collection (56 DAS).

Trichoderma harzianum had a higher effectiveness compared to the control treatment on the number of branches and leaves. Table 8 shows *Trichoderma harzianum*, for instance, in trial one increased the number of branches from 2.00 at 14 DAS to 4.25 at 28 DAS to 6.58 at 42 DAS to 7.92 at 56 DAS, compared to the control, which increased from 1.58 at 14 DAS to 2.67 at 28 DAS to 3.92 at 42 DAS to 4.50 at 56 DAS. In trial two, Table 9 shows *Trichoderma harzianum* increased the number of leaves from 4.75 at 14 DAS to 12.25 at 28 DAS to 30.00 at 42 DAS to 39.25 at 56

DAS compared to the control, which increased from 3.50 at 14 DAS to 8.75 at 28 DAS to 18.25 at 42 DAS to 23.00^d at 56 DAS. *Trichoderma harzianum* had a lower effect on shoot development compared to the direct application of nutrient amendments like NPK. Control treatment was the last in terms of the number of branches and leaves compared to all other treatments. This is because there were no essential nutrients added to the soil to promote plant growth.

These findings are in line with those of Raksun *et al.* (2022), who reported that NPK fertilizer increased the number of leaves, leaf width, stem length, and leaf length of long beans. These findings are consistent with those of Ali *et al.* (2023), who reported that organic fertilizers applied at the rate of 600 kg ha⁻¹ increased the number of branches compared to the control in common beans. These findings corroborate with those of Sofo *et al.* (2010), who reported that there was increased root growth and reduced shoot growth in prunus rootstock. These results are inconsistent with those of El-Yazal (2020), who reported that FYM increased the number of branches compared to NPK and control treatment in broad beans. These findings are contrary to those of Naluzze (2022), who reported that there was no significant difference ($P>0.05$) in the number of leaves when NPK and cattle manure were applied compared to the control in common beans.

4.2.4 Effect of NPK (23.23.0), Farmyard manure, and *Trichoderma harizianum* on the Number of Pods and Seeds of Common Beans

Table 10 represents the number of common bean pods and seeds produced, respectively. There was a significant difference ($p<0.05$) (Appendix 59 to 63) in all the treatments in both trials in terms of the number of pods and seeds. The NPK treatment was the most effective treatment with the highest number of pods and seeds in both trials compared to the control treatment. There were similar trends observed in both trials. As presented in Table 10, looking at the trends in both trials on the number of pods and seeds, NPK had the highest, followed by FYM, then *Trichoderma harzianum*, and Control was the last. The NPK treatment supplied readily available nutrients, which accelerated growth, hence the highest number of pods and leaves, while FYM slowly released nutrients for common bean uptake, improved water retention, soil structure, and microbial activity for plant growth. *Trichoderma harzianum* enhanced nutrient uptake but at a lesser extent compared to NPK and

FYM, root health indirectly supporting moderate growth. In comparison to the control, which depended solely on existing soil fertility, which limited common bean development and reduced overall growth performance.

Table 10: Effect of NPK (23.23.0), Farmyard manure, and *Trichoderma harizianum* on the Number of Pods and Seeds of Common Beans.

| TRIAL | TREATMENT | NUMBER OF PODS | NUMBER OF SEEDS |
|-------|-----------------------|--------------------|---------------------|
| 1 | Control | 6.00 ^d | 33.70 ^d |
| | NPK | 25.10 ^a | 176.77 ^a |
| | Farmyard manure | 17.37 ^b | 121.63 ^b |
| | TH | 11.33 ^c | 64.90 ^c |
| 2 | Control | 5.73 ^d | 31.63 ^d |
| | NPK | 25.57 ^a | 162.00 ^a |
| | Farmyard manure | 18.03 ^b | 112.13 ^b |
| | TH | 11.37 ^c | 69.77 ^c |
| | Mean | 15.06 | 96.57 |
| | CV (%) | 20.01 | 26.51 |
| | LSD _(0.05) | 1.085 | 9.21 |

Legend: TH- *Trichoderma harzianum*, CV- Coefficient of variation, Means with the same letters within the column are not significantly different at (P <0.05).

As shown in Table 10 NPK treatment had the highest number of pods 25.10 (trial one) and 25.57 (trial two) and seeds 176.77 (trial one) and 162.00 (trial two) compared to control which had the least number of pods 6.00 (trial one) and 5.73 (trial two) and seeds 33.70 (trial one) and 31.63 (trial two). The NPK treatment emerged as the best because of the supply of essential macronutrients, which played vital roles in flowering, pod development, seed formation, and filling. Farmyard manure had the second highest treatment number of pods of 17.37 (trial one) and 18.03 (trial two) and number of seeds were 121.63 (trial one) and 112.13 (trial two) compared to control which had the least number of pods 6.00 (trial one) and 5.73 (trial two) and seeds 33.70 (trial one) and 31.63 (trial two) in both trials. Farmyard manure improves the soil's physical properties and aeration, and enhances water retention and infiltration. It aids in the slow release of micro and macro nutrients, which become available to the common bean plant. Moreover, it enriches microbial populations in soil, leading to a healthy rhizosphere that can help in nutrient uptake and nutrient cycling. Farmyard manure has long-term benefits like sustained fertility and enhanced soil structure, which indirectly contribute to an increased number of pods and seeds.

Trichoderma harzianum had 11.33 (trial one) with and 11.37 (trial two) number of pods and seeds with 64.90 (trial one) and 69.77 (trial two) compared to control treatment which had the lowest number of pods 6.00 (trial one) and 5.73 (trial two) and seeds 33.70 (trial one) and 31.63 (trial two) in both trials. *Trichoderma harzianum* is known to promote growth through root development, disease suppression, and nutrient solubilisation, but to a lesser extent compared to NPK and FYM. Control treatment had the lowest number of pods of 6.00 (trial one) and 5.73 (trial two), and seeds 33.70 (trial one) and 31.63 (trial two) in both trials. This shows that there is a need for intervention in order to increase the production of seeds and pods. These findings corroborate with those of Wijanarko *et al.* (2016), who reported that the application of NPK fertilizer increased the number of pods compared to the control in soybean.

Similar findings were reported by Zamukulu *et al.* (2023), who found that NPK fertilizer increased the number of seeds per plant compared to the control. These findings aligned with those of Tadesse & Abera (2023), who reported that application of farmyard manure increased the number of pods and seeds per plant in haricot bean. These findings align with those of Ali *et al.* (2023), who reported that organic fertilizers applied at the rate of 600 kg ha⁻¹ increased the number of seeds and number of pods compared to the control in common beans. These findings corroborate with those of Mahmoodian *et al.* (2022), who reported that *Trichoderma harzianum* increased the number of pods and seeds compared to the control. These results are inconsistent with those of El-Yazal (2020), who reported that FYM increased the number of seeds and branches compared to NPK and control treatment in broad beans. These findings are contrary to those of Naluzze (2022), who reported that there was no significant difference ($P > 0.05$) in the number of pods and seeds when NPK and cattle manure were applied compared to the control in common beans.

4.2.5 Effect of NPK (23.23.0), Farmyard Manure, and *Trichoderma harizianum* on Grain and Biomass Yields of Common Beans

Biomass produced is an essential indicator of plant vigour and the ability to capture as well as convert sunlight energy to usable energy by the process of photosynthesis, which is linked to water-use-efficiency and nutrient uptake. As presented in Table 11, there were significant differences ($p < 0.05$) (Appendix 65 to 69) in grain and biomass

yields of common beans among NPK, FYM, and *Trichoderma harzianum* treatments. Still, *Trichoderma harzianum* and control did not show a significant difference ($p>0.05$) in yields (trials 1 and 2) and biomass (trial 1). In trial two, there were significant differences ($p<0.05$) among all treatments in biomass.

Table 11: Effect of NPK (23.23.0), Farmyard manure, and *Trichoderma harizianum* on Grain and Biomass Yields of Common Beans.

| TRIAL | TREATMENT | GRAIN YIELDS (kg/ha) | BIOMASS (kg/ha) |
|-------|------------------------------|----------------------|----------------------|
| 1 | Control | 385.19 ^c | 196.88 ^c |
| | NPK | 3950.62 ^a | 1269.25 ^a |
| | Farmyard manure | 2437.67 ^b | 656.53 ^b |
| | <i>Trichoderma harzianum</i> | 923.33 ^c | 372.27 ^c |
| 2 | Control | 787.65 ^c | 189.59 ^d |
| | NPK | 5234.57 ^a | 1154.18 ^a |
| | Farmyard manure | 3061.73 ^b | 714.67 ^b |
| | <i>Trichoderma harzianum</i> | 1354.32 ^c | 361.01 ^c |
| | Mean | 2192.81 | 614.30 |
| | CV (%) | 31.85 | 13.25 |
| | LSD _(0.05) | 850.76 | 99.12 |

Legend: CV- Coefficient of variation; Means with the same letters within the column are not significantly different at ($P < 0.05$)

As shown in Table 11, looking at the trends in both trials on the increase of grain yield and biomass, NPK had the highest, followed by FYM, then *Trichoderma harzianum*, and Control had the least. The NPK treatment supplied readily available nutrients, which accelerated growth, hence the highest grain yield and biomass, while FYM slowly released nutrients for common bean uptake, improved water retention, soil structure, and microbial activity for plant growth. *Trichoderma harzianum* enhanced nutrient uptake but at a lesser extend compared to NPK and FYM, root health indirectly supporting moderate growth. In comparison to the control, which depended solely on existing soil fertility, which limited common bean development and reduced overall growth performance. The NPK treatment had the highest grain yields of 3950.62 (trial 1) and 5234.57 (trial 2), and biomass of 1269.25 (trial 1) and 1154.18 (trial 2), while control had the lowest yield of 385.19 (trial1) and 787.65 (trial 2) and biomass of 196.88 (trial1) and 189.59 (trial two) in both trials.

The NPK treatment provided readily available nutrients for physiological processes. The performance was contributed by readily available nutrients, i.e., nitrogen and phosphorus, which are essential for leaf development, cell division, energy transfer,

and root expansion within common beans. The results showed the effectiveness of NPK fertilizer in maximising common bean growth and biomass, which also contributed to increased yield. Farmyard manure was the second among other treatments with yield of 2437.67 (trial 1) and 3061.73 (trial 2) and biomass of 656.53 (trial 1) and 714.67 (trial 2) compared to control which had the lowest yield of 385.19 (trial1) and 787.65 (trial 2) and biomass of 196.88 (trial1) and 189.59 (trial two) in both trials. Farmyard manure showed the value of organic inputs in improving crop productivity and soil fertility, because of enhanced water retention and soil structure. Though it was less effective compared to NPK, farmyard manure offers agronomic and environmental advantages. The increase in the amount of biomass under farmyard manure application shows its ability to become an organic alternative to synthetic fertilizers.

Trichoderma harzianum as a bio-fertilizer illustrated a grain yield of 923.33 (trial 1) and 1354.32 (trial 2) and biomass of 372.27 (trial 1) and 361.01 (trial 2) in comparison to control, which had the lowest grain yield of 385.19 (trial 1) and 787.65 (trial 2) and biomass of 196.88 (trial 1) and 189.59 (trial two) in both trials. However, the grain yield of *Trichoderma harzianum* was lower compared to FYM and NPK. It showed its potential in increasing vegetative growth by nutrient mobilisation and enhanced root health. Control treatment had the lowest grain yield of 385.19 (trial 1) and 787.65 (trial 2), and biomass of 196.88 (trial 1) and 189.59 (trial 2) produced in both trials, but there was no statistically significant difference ($p>0.05$) with *Trichoderma harzianum* except on biomass (trial 2). This reflected that there was a nutrient deficiency in the soil, and this suggests external inputs should be added to the soil in order to support vegetative growth.

These findings, which corroborate with those of Mndzebele *et al.* (2023), showed that application of NPK fertilizer increased biomass accumulation in cowpea-amaranth intercrop. These findings align with those of Zamukulu *et al.* (2023), who observed that application of 150 kg per ha of NPK fertilizer led to increased yield of common bean from 1.1 to 1.5 tons per ha. These results are consistent with those of Rurangwa *et al.* (2018), who demonstrated that manure significantly increased grain yield and biomass of soybean and common bean in comparison to untreated soils. These results

corroborate with those of Upenji (2020), who showed that manure improved the yield of common bean compared to the control.

These findings corroborate with those of Ali *et al.* (2023), who reported that organic fertilizers applied at the rate of 600 kg ha^{-1} increased the grain yield and dry matter compared to the control in common beans. These findings are in line with those of Pereira *et al.* (2014), who reported that *Trichoderma harzianum* improved the growth of common bean; hence, there is a potential increase in yield. These results comply with those of Mahmoodian *et al.* (2022), who reported that inoculation of *Trichoderma harzianum* led to increased total biomass compared to the control. These results are inconsistent with those of El-Yazal (2020), who reported that FYM increased yields compared to NPK and control treatment in broad beans. These findings are contrary to those of Alhrout *et al.* (2018), who reported that FYM showed increased yield compared to NPK and control in tomato.

4.3 Effect of NPK (23.23.0), *Trichoderma Harzianum*, and Farmyard Manure on Plant-parasitic Nematodes in the Roots of Common Beans.

4.3.1 Effect of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on the Number of Common Beans Root Nodules

Table 12 presents the number of nodules from two trials determining the effect of different treatments on the number of nodules of common beans. The treatments were; *Trichoderma harzianum*, NPK, control, and farmyard manure. In every trial, the average number of nodules was reported for each treatment, together with a statistical comparison indicated by the letters a, b, c, and d. The table also provides an overall "Mean" number of nodules across all treatments and trials, a "CV (%)" (Coefficient of Variation) indicating the variability of the data, and "LSD" (Least Significant Difference at the 0.05 level), which was used to determine statistically significant differences among treatment means.

Table 12: Effect of NPK (23.23.0), Farmyard manure, and *Trichoderma harzianum* on the Number of Root nodules of Common Beans.

| TRIAL | TREATMENT | NUMBER OF NODULES |
|-------|------------------------------|--------------------|
| 1 | Control | 5.56 ^c |
| | NPK | 2.44 ^c |
| | Farmyard manure | 14.11 ^b |
| | <i>Trichoderma harzianum</i> | 41.00 ^a |
| 2 | Control | 17.67 ^c |
| | NPK | 3.44 ^d |
| | Farmyard manure | 32.89 ^b |
| | <i>Trichoderma harzianum</i> | 60.00 ^a |
| | Mean | 22.14 |
| | CV (%) | 34.61 |
| | LSD _(0.05) | 5.102 |

Legend: CV- Coefficient of variation, Means with the same letters within the column are not significantly different at (P <0.05)

Nodulation in common beans is an essential parameter that shows the symbiotic interaction between nitrogen-fixing bacteria and the plant. Soil amendments like organic inputs, bioinoculants, and inorganic fertilizers influence these symbiosis interactions. Table 12 shows that there were significant differences ($p < 0.05$) (Appendix 71 and 72) among the treatments in terms of their effect on the nodulation of common beans. Looking at the trends in both trials on the number of nodules, *Trichoderma harzianum* had the highest, as shown in Figure 5, followed by FYM, as shown in Figure 2, then Control, as shown in Figure 3, and NPK, as shown in Figure 4, had the least. *Trichoderma harzianum* increased nodules by encouraging symbiosis with rhizobia and root health, resulting in more nodules. Farmyard manure improved microbial activity and soil fertility, hence boosting nodulation. The NPK treatment input reduced rhizobial infection, leading to least nodules, while the control had no amendment added.

Trichoderma harzianum had the highest number of nodules in both trials, with 41.00 in trial one and 60.00 in trial two, and NPK had the lowest number, 2.44 in trial one and 3.44 in trial 2, in comparison to the control, which had the fewest nodules, 5.56 in trial one and 17.67 in trial 2. *Trichoderma harzianum* promoted nodulation in common beans by enhancing root growth and microbial activity, hence improved rhizobial colonisation. *Trichoderma* secretes metabolites like indole-3-acetic acid, which promote rhizobia interaction. Farmyard manure was the second in the number of nodules in both trials, with 14.11 in trial 1 and 32.89 in trial two, compared to the

control, which had 5.56 in trial one and 17.67 in trial 2, and its effects were less compared to *Trichoderma harzianum*, since it does not have directly bioactive compounds which stimulate nodulation. Farmyard manure can enhance soil structure and improve microbial biomass, and nutrients are released slowly, which favours the common bean plant and rhizobia.

As shown in Figure 4, NPK had the lowest number of nodules. The NPK treatment had the lowest number of nodules of 2.44 in trial one and 3.44 in trial 2, compared to the control, which had 5.56 in trial one and 17.67 in trial 2. The NPK treatment did not support the growth of root nodules since it presented a suppressive effect in terms of nodulation. The control treatment had a higher number of nodules compared to NPK, showing that nodulation occurred naturally in the soil but at a slower rate than in FYM and *Trichoderma harzianum* treatments. These findings corroborate with those of Mweetwa *et al.* (2016), who reported that inoculation of *Trichoderma* led to an increased number of nodules.

These findings are consistent with those of Otieno *et al.* (2009), who carried out a study involving legumes like lablab, lima bean, green gram, and common bean, and found that application of farmyard manure increased the number of root nodules compared to the control treatment. This outcome was contributed to by slow mineralisation, resulting in the gradual release of nitrogen as well as phosphorus in the farmyard manure. These findings are in line with those of Dos *et al.* (2022), who reported that application of nitrogen fertilizer reduced the reliance of the plant on biological nitrogen fixation. When mineral nitrogen is available, legumes do not concentrate on nodules formation since they can get the nitrogen required by the crop without involving symbiosis. These findings are contrary to those of Sulieman & Hago (2009), who reported that farmyard manure did not have a significant effect on nodulation compared to control in common beans.



Figure 2: Common bean root from Farmyard manure treatment



Figure 3: Common bean root from control treatment



Figure 4: Common bean root from NPK (23.23.0) treatment



Figure 5: Common bean root from *Trichoderma harzianum* treatment

4.3.2 Effect of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard Manure on Plant-parasitic Nematodes in Roots of Common Bean

Table 13 shows results from trial one and trial two, determining the effect of different treatments (farmyard manure, *Trichoderma harzianum*, NPK, and control) on PPNs populations in the roots of common beans. The PPNs identified were: *Tylenchus spp*, *Helicotylenchus spp*, *Rotylenchus spp*, and *Pratylenchus spp*.

Table 13: The Number of Plant-parasitic Nematodes Identified in Roots (5g) of Common Beans under NPK, *Trichoderma harzianum*, and Farmyard Manure

| Trial | TRT | <i>Tylenchus spp</i> | <i>Helicotylenchus spp</i> | <i>Rotylenchus spp</i> | <i>Pratylenchus spp</i> | total |
|-------|---------|----------------------|----------------------------|------------------------|-------------------------|-------|
| 1 | Control | 0 | 18 | 0 | 0 | 18 |
| | NPK | 4 | 19 | 10 | 10 | 43 |
| | FYM | 0 | 0 | 0 | 0 | 0 |
| | TH | 0 | 0 | 0 | 0 | 0 |
| 2 | Control | 16 | 5 | 12 | 30 | 63 |
| | NPK | 0 | 12 | 7 | 30 | 49 |
| | FYM | 0 | 0 | 0 | 10 | 10 |
| | TH | 5 | 0 | 0 | 0 | 5 |

Legend: TH. - *Trichoderma harzianum*; FYM- Farmyard manure; TRT- Treatment

Table 14: Effect of NPK, *Trichoderma harzianum*, and Farmyard Manure on the Plant-parasitic Nematodes in Roots (5g) of Common Beans

| TRIAL | TREATMENT | PPNs in ROOTS |
|-------|-----------------------|-------------------|
| 1 | Control | 18 ^{ab} |
| | NPK | 43 ^a |
| | FYM | 0.14 ^b |
| | TH | 0.00 ^b |
| 2 | Control | 63 ^a |
| | NPK | 49 ^a |
| | FYM | 10 ^b |
| | TH | 5 ^b |
| | Mean | 23.56 |
| | CV (%) | 54.28 |
| | LSD _(0.05) | 19.02 |

Legend: TH - *Trichoderma harzianum*; FYM- Farmyard manure; TRT- Treatment
Means with the same letters within the column are not significantly different at ($P < 0.05$).

As shown in Table 14 in trial one, control (43) was significantly different ($p < 0.05$) from FYM (0.14) and TH (0.00). NPK (18) was not significantly different ($p > 0.05$) (Appendix 74) from the control. *Trichoderma harzianum* and FYM significantly

reduced plant-parasitic nematodes, showing suppression of PPNs in comparison to the control. In trial two, control (63) and NPK (49) were similar statistically but showed significant different ($p < 0.05$) (Appendix 75) from FYM (10) and TH (5). Control and NPK caused an increase in the number of PPNs, while *Trichoderma harzianum* and FYM reduced the numbers. The control treatment increased PPNs in the roots naturally because there were no amendments added to suppress PPNs. The NPK treatment encouraged plant growth, especially vigorous root growth, which provided feeding sites for PPNs, which increased the population of PPNs. Farmyard manure reduced plant-parasitic nematodes by encouraging free-living nematodes that suppress PPNs. *Trichoderma harzianum* suppressed PPNs through competition, parasitism, and producing nematicidal metabolites.

As presented in Table 13 in trial 1, NPK treatment had the highest number of PPNs (43) compared to control, which had 18, but FYM and *Trichoderma harzianum* did not have PPNs. In trial two, control treatment had the highest number of plant-parasitic nematodes (63), followed by NPK (49), then FYM (10), and *Trichoderma harzianum* treatment had (5). Farmyard manure and *Trichoderma harzianum* showed to have suppressed PPNs. Control treatment had the highest number of PPNs, especially in trial two, because there was no biotic suppression mechanism. For instance, in trial two, control treatment had *Tylenchus spp* (16), *Helicotylenchus spp* (5), *Rotylenchulus spp* (12), and *Pratylenchus spp* (30) compared to FYM, which had *Pratylenchus spp* (10), *Trichoderma harzianum* had *Tylenchus spp* (5). Control treatment made the roots of common beans highly susceptible to PPNs invasion. When nutrients are not added or microbial communities are not present, the defence mechanism is weakened, allowing plant-parasitic nematodes like *Tylenchus spp*, *Helicotylenchus spp*, *Rotylenchulus spp*, and *Pratylenchus spp* to thrive freely in the vascular tissues and root cortex (Abd-Elgawad, 2020). PPNs can result in lesions and root galling, hence interfering with the root function.

In trial two, NPK treatment was the second in terms of the number of PPNs. The NPK treatment increased PPNs in roots since it provided readily available nutrients for common bean growth, hence providing feeding sites and a conducive habitat for PPNs. Farmyard manure reduced the number of PPNs compared to the control treatment. For example, in trial one, the control treatment had *Helicotylenchus spp*

(18) compared to FYM, which did not have PPNs. This illustrated the role of FYM in protecting common bean roots. Farmyard manure improved the condition of the root zone by increasing populations of microbial antagonists, increasing root biomass, and creating an uncondusive environment for egg hatching and PPNs' survival. *Trichoderma harzianum* had the lowest count of PPNs, with no PPNs in trial one and (5) in trial two, compared to the control, which had (18) in trial one and (63) in trial 2. *Trichoderma harzianum* colonized cortex tissues and root surfaces, producing enzymes and secondary metabolites that degrade PPN eggs as well as inhibit penetration by the juvenile. Moreover, *Trichoderma harzianum* competed for root space, hence making a protective barrier against invasion by nematodes.

These findings are consistent with those of Neher *et al.* (2010), who reported that *Trichoderma harzianum* also promoted healthier common bean root systems that tolerated infections. These findings are consistent with those of Deepali (2023), who reported that synthetic fertilizers provide feeding sites for PPNs. These findings are in line with those of Pinto *et al.* (2024), who determined that *Trichoderma sp* protects the roots from PPNs and also influences nematodes, i.e, fungivores. These findings comply with those of Kimenju *et al.* (2009), who reported that FYM also created microbial competition in the roots, hence leading to PPN suppression. In contrast to these findings, Lawal & Atungwu (2017) reported that NPK fertilizer increased plant-parasitic nematodes compared to the control treatment, while organic fertilizer suppressed PPNs.

CHAPTER FIVE

SUMMARY, CONCLUSION AND RECOMMENDATIONS

5.1 Summary of Findings

Objective one sought to describe the effects of NPK (23.23.0), *Trichoderma harzianum*, and Farmyard manure on Numbers, Evenness, and Species Diversity of Soil Nematodes under Common Beans. Looking at the number of plant-parasitic and free-living nematodes that were identified, *Trichoderma harzianum*, and Farmyard manure increased FLN. For example, in trial two, *Trichoderma harzianum* increased FLN from 406 (initial) to 412 (flowering stage) to 745 (final) while Farmyard manure increased from 456 (initial) to 831 (flowering stage) to 1426 (final), in comparison to control and NPK, which reduced the number of FLN. For instance, in trial one, the control treatment reduced the number of FLN from 615 (initial) to 277 (flowering stage) to 156 (final), and NPK reduced from 459 (initial) to 188 (flowering stage) to 133 (final). Control and NPK led to an increase in plant-parasitic nematodes. For instance, in trial two, control increased PPNs from 477 (initial) to 520 (flowering stage) to 1068 (final), and NPK increased from 368 (initial) to 380 (flowering stage) to 724 (final). However, *Trichoderma harzianum* and Farmyard manure reduced the number of PPNs, which showed their suppressive effect on these nematodes. For instance, in trial two, FYM reduced the number of PPNs from 296 (initial) to 115 (flowering stage) to 48 (final), and *Trichoderma harzianum* reduced from 237 (initial) to 53 (flowering) to 9 (final).

Species diversity of soil nematodes was calculated using Shannon index of diversity and Simpson's index of diversity, while Pielou's index determined Evenness. *Trichoderma harzianum* increased the Shannon index of diversity of FLN, for instance, in trial one from 1.16 (initial) to 1.24 (final). In comparison, the control treatment reduced the Shannon index of diversity of FLN from 1.26 to 0.79 in trial one. *Trichoderma harzianum* increased the Simpson's diversity index of FLN from 0.67 (initial) to 0.73 (final) while the control reduced diversity from 0.53 (initial) to 0.54 (final) in trial two. *Trichoderma harzianum* had the highest Evenness of 0.91 in trial two, while the control had 0.82. Control and NPK increased the Shannon index of diversity and Simpson's index of diversity of PPNs. For instance, in trial one, control increased Shannon diversity of PPNs from 1.04 (initial) to 1.71 (final), and NPK increased from 0.99 (initial) to 1.41 (final). For example, in trial one, control

increased Simpson's index of diversity from 0.56 (initial) to 0.78 (final), and NPK increased from 0.61 (initial) to 0.70 (final) while *Trichoderma harzianum* reduced from 0.66 (initial) to 0.25 (final). *Trichoderma harzianum* increased the Evenness of PPNs from 0.73 (initial) to 0.96 (final).

Objective two sought to describe the effect of NPK (23.23.0), *Trichoderma harzianum*, and farmyard manure on common beans' growth and yield components. Plant parameters which were measured were height, number of leaves, and number of leaves. The yield components were the number of pods, seeds, and yield. Biomass was also measured after harvesting beans. The NPK treatment had the highest plant height, number of branches, and number of leaves while the control had the least across the growth period. For example, in trial two at 14 days after sowing, NPK had 16.25^a cm, FYM 13.25^b cm, *Trichoderma harzianum* 10.58^c cm, and control 8.08^d cm after conducting LSD, where alpha was 0.05. The results showed that there was a significant difference ($p < 0.05$) between the treatments. The NPK treatment also increased the number of pods, seeds, yield, and biomass. In trial one, the number of pods, NPK had 25.10^a, FYM 17.37^b, *Trichoderma harzianum* 11.33^c and control 6^d. Number of seeds: NPK had 176.77^a, FYM 121.63^b, *Trichoderma harzianum* 64.90^c and control 33.7^d. For grain yield: NPK had 3950.62^a kg/ha, FYM 2437.67^b kg/ha, *Trichoderma harzianum* 923.33^c kg/ha, and control 385.19^c kg/ha. Biomass; NPK had 1154.18^a kg/ha, FYM 714.67^b kg/ha, *Trichoderma harzianum* 361.01^c kg/ha, and control 189.59^d kg/ha. These results showed that there was a significant difference ($p < 0.05$) between treatments.

Objective three described the effect of NPK (23.23.0), *Trichoderma harzianum*, and farmyard manure on plant-parasitic nematodes in the roots of common beans. The numbers of nodules in the roots of common beans were counted, with *Trichoderma harzianum* having the highest and NPK recording the lowest number of nodules. In trial two, *Trichoderma harzianum* had 60^a, FYM 32.89^b, control 17.67^c and NPK 3.44^d. These results showed that there were significant differences ($p < 0.05$) among the treatments. Plant-parasitic nematodes in the roots were identified and counted, where control and NPK increased the number of PPNs, while *Trichoderma harzianum* and FYM reduced their numbers. For instance, in trial one, NPK had the highest

number of PPNs in the roots (43), while FYM and *Trichoderma harzianum* treatments did not have PPNs in the roots.

5.2 Conclusion

In conclusion, *Trichoderma harzianum* and FYM increased FLN abundance (Table 2) and reduced PPNs in the soil (Table 4). Farmyard manure and *Trichoderma harzianum* resulted in the highest overall nematode abundance of FLN, particularly increasing populations of *Dorylaimus* and *Bacterivores* (Table 1), indicating improved soil biological activity. Control and NPK increased the number of PPNs (Table 4) and reduced the FLN in soil (Table 2). *Trichoderma harzianum* applications increased free-living nematode diversity (Table 5) while stabilising or reducing plant-parasitic nematode diversity (Table 5). Control treatment increased the diversity of PPNs (Table 5) and decreased that of FLN (Table 5). The NPK treatment exhibited the highest values for plant parameters (Tables 7, 8, 9), yield components (Tables 10 and 11), and biomass (Table 11), with significant differences ($p < 0.05$) from the other treatments. In contrast, the control had the lowest values for all these parameters. *Trichoderma harzianum* and FYM reduced PPNs in roots of common beans, while control and NPK increased the abundance of PPNs in roots (Table 14). *Trichoderma harzianum* had the highest number of nodules and showed a significant difference ($p < 0.05$) from other treatments (Table 12). Overall, these results confirm that organic and biological amendments enhance soil nematode diversity, fertility, and productivity, contributing to sustainable bean cultivation.

5.3 Recommendation

The study recommends the following:

- i. Integration of *Trichoderma harzianum* as a biological amendment to promote soil biodiversity and suppress plant-parasitic nematodes through natural competition and antagonism. Incorporating farmyard manure is highly recommended for promoting soil biological activity, enhancing nematode diversity, and improving soil health. Combining *Trichoderma harzianum* with organic amendments like FYM could balance soil fertility while enhancing biological control agents. Reliance on chemical fertilizers, such as NPK, should be reduced to prevent the suppression of free-living nematodes.

- ii. Application of *Trichoderma harzianum* and FYM promotes soil health and increases growth and yield of common beans. Long-term use of FYM is the better option for soil health and yields. Low yields recorded by the control treatment showed that external inputs should be added to the soil.
- iii. Continuous assessment of nematode communities over multiple growing seasons is essential to understand the long-term impacts of different soil amendments on diversity and crop productivity. Application of *Trichoderma harzianum* is recommended as it enhances root nodulation in common beans, thereby improving nitrogen fixation.

5.4 Suggestion for Further Studies

Further studies can be done on soil amendments that increase free-living nematodes, since they are able to suppress plant-parasitic nematodes and promote soil health and crop productivity.

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APPENDICES

Appendix 1: Identification Key for Major Plant-Parasitic Nematodes As described in Mekete *et al.*, (2012)

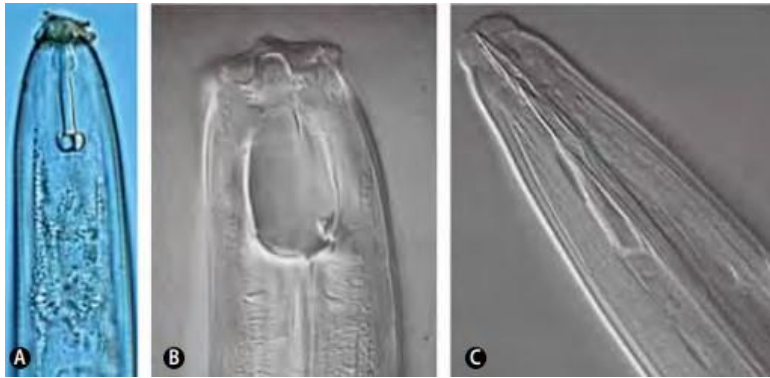


Figure 1: Stoma with stylet (A) and without (B,C)

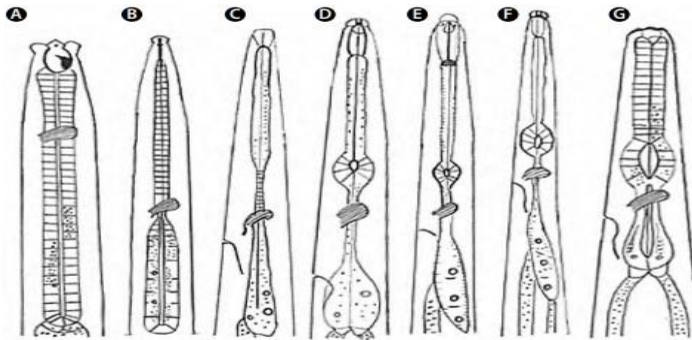


Figure 2: Different types of esophagi: one part (A), two part Dorylaimoid (B), three part (C), four part TYlenchoid (D,E,F) and four part Rhabditoid (G)

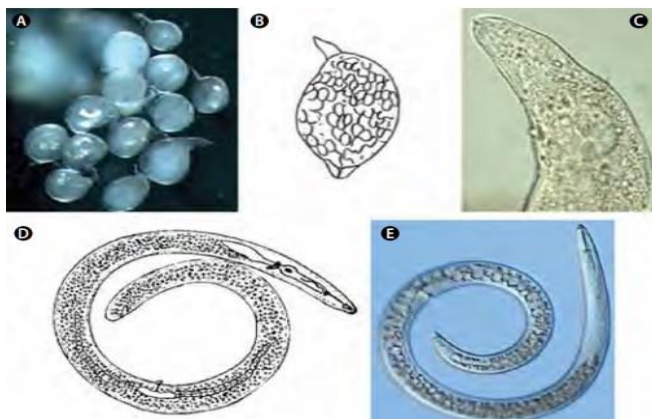


Figure 3: Female nematode body types, swollen (A, B, C) and cylindrical (D, E)

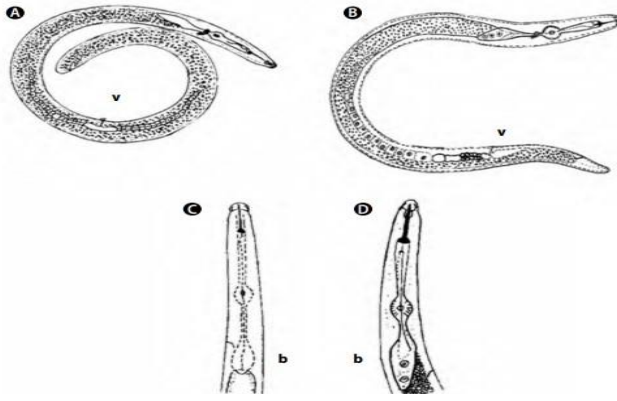


Figure 4: (A) Vulva [v] location near mid body, (B) posteriorly toward tail, (C) basal bulb [b] without overlap, and (D) intestines with overlap.

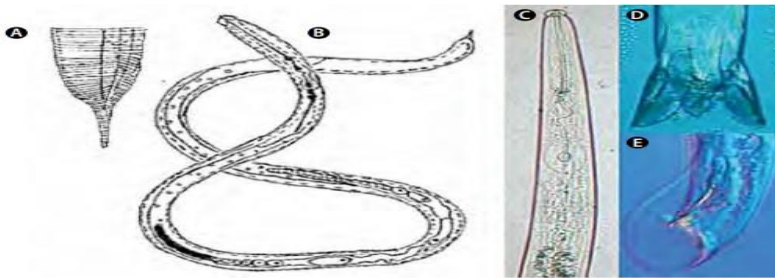


Figure 5: Dolichodoros female and juvenile tail shape (A), female full body (B), head region (C), male tail dorsal view (D), and male tail lateral view (E).

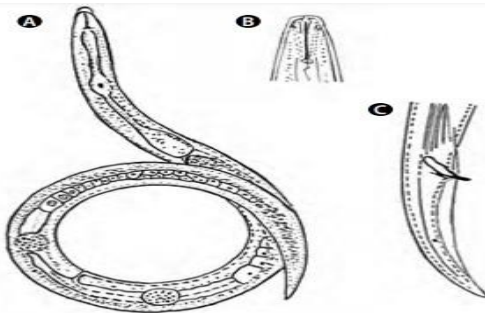


Figure 6: Merlinius female (A), stylet region (B), and male tail (C).

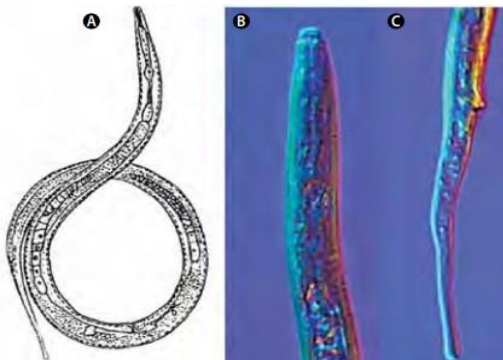


Figure 7: Psilenchus full body (A), head region (B), and male tail (C).

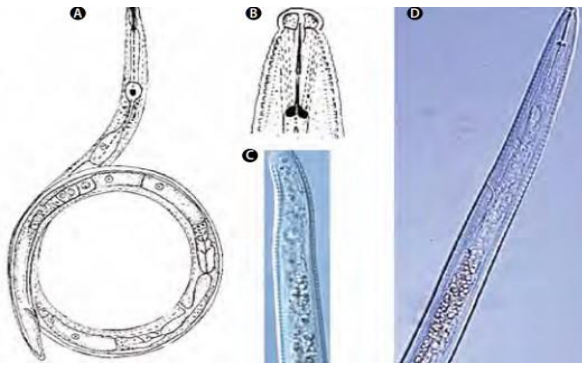


Figure 8: *Tylenchorhynchus* full body (A), short stylet (B), round tail tip (C), basal bulb not overlapping intestine (D).

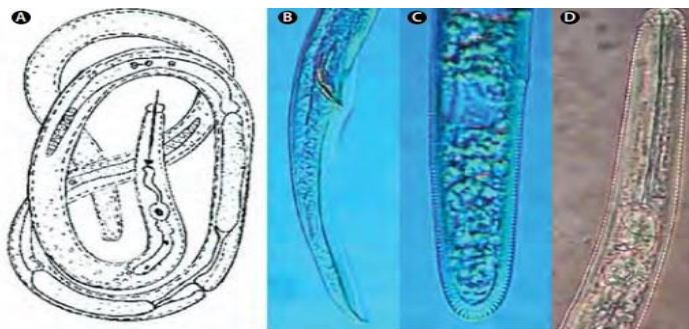


Figure 9: *Belonolaimus* full body length (A), male tail region (B), female tail region (C), and head region (D).

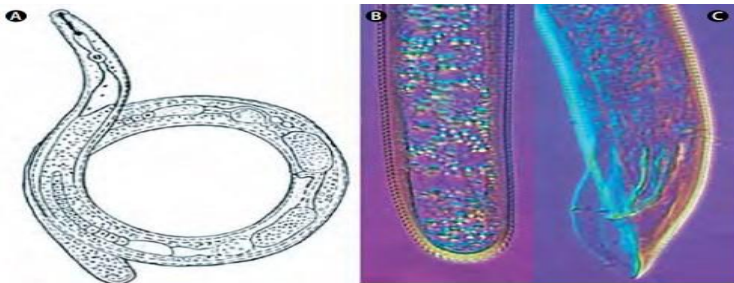


Figure 10: *Hoplolaimus* full body (A), female tail (B), and male tail (C).

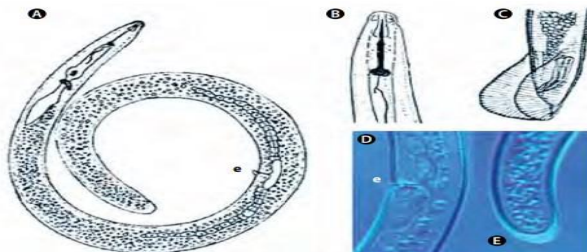


Figure 11: *Peltamigratus* female full body with epitygma [e] (A), stylet region (B), male tail (C), epitygma (D), and female tail region (E).

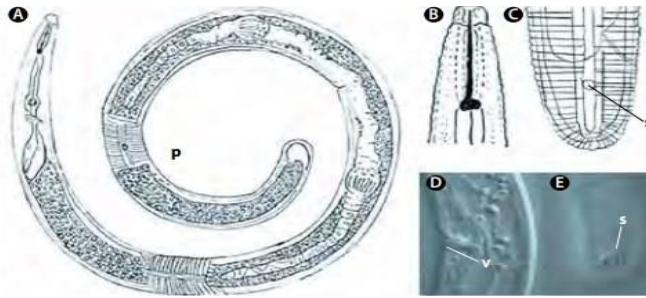


Figure 12. Aorolaimus female body with phasmid [p] midbody (A), stylet region of Aorolaimus and Scutellonema (B), tail region of Scutellonema with scutellum [s] (C), vulva [v] without epitygma (D), and scutellum [s] near tail tip of Scutellonema (E).

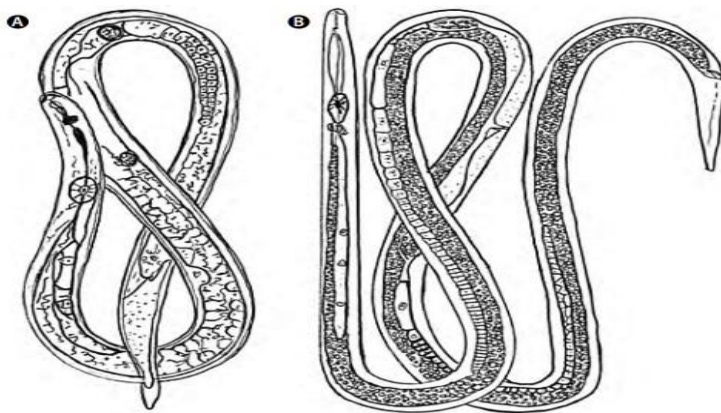


Figure 13. Female full body of Radopholus (A) and Hirschmanniella (B).

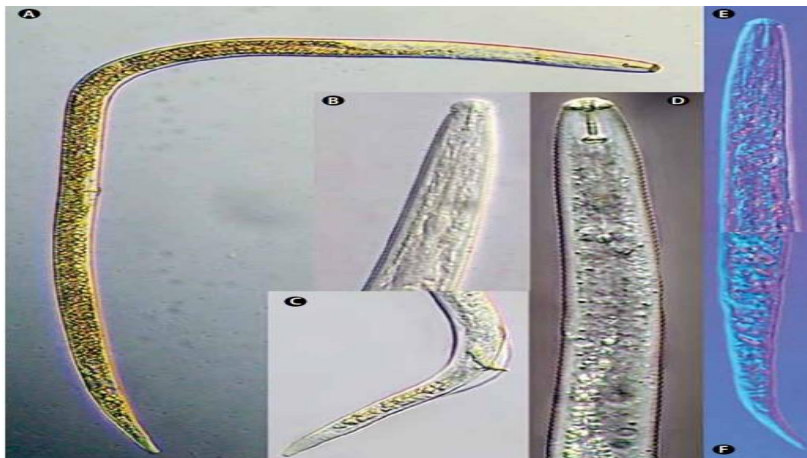


Figure 14. Radopholus mature female full body (A), male anterior head (B), male tail (C), female anterior end (D); Hirschmanniella anterior end (E) and tail region (F).

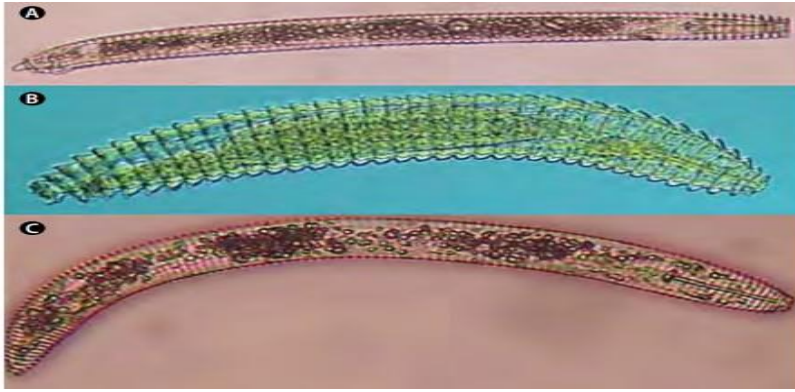


Figure 15. Entire body of Hemicriconemoides (A), Criconema (B), and Criconemella (C).

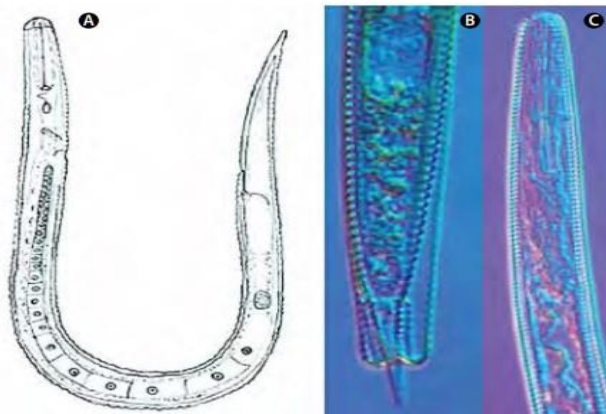


Figure 16. Hemicycliophora female full body (A), tail (B), and head (C).

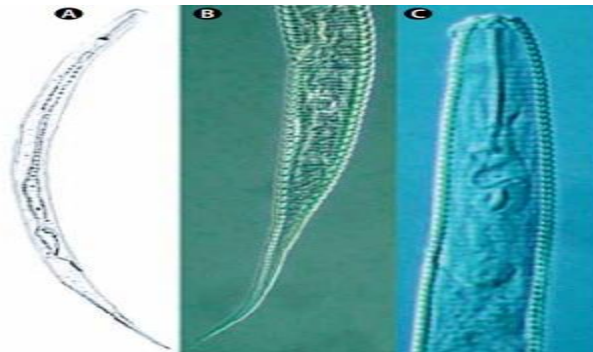


Figure 17. Caloosia female body (A), tail (B), and head (C).

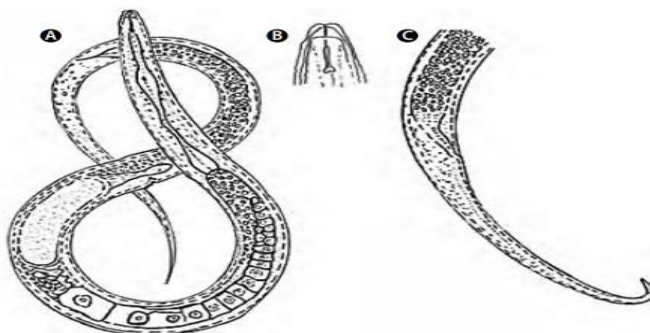


Figure 18. Tylenchulus full body view (A), stylet region (B), and tail (C).

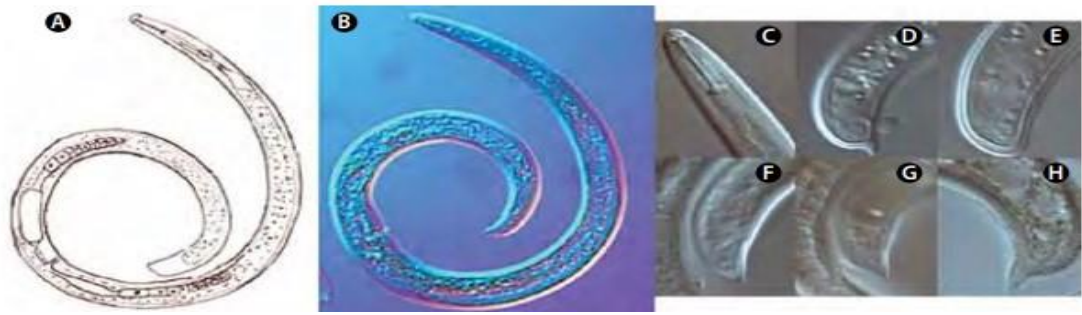


Figure 19. Full body spiral habitus of *Helicotylenchus* spp., (A, B), dorsal gland opening distance from stylet end (C), rounded tail with terminal projection (D), hemispherical annulated tail terminus (E), irregular tail projections (F, G), tail with non-annulated ventral projection (H).

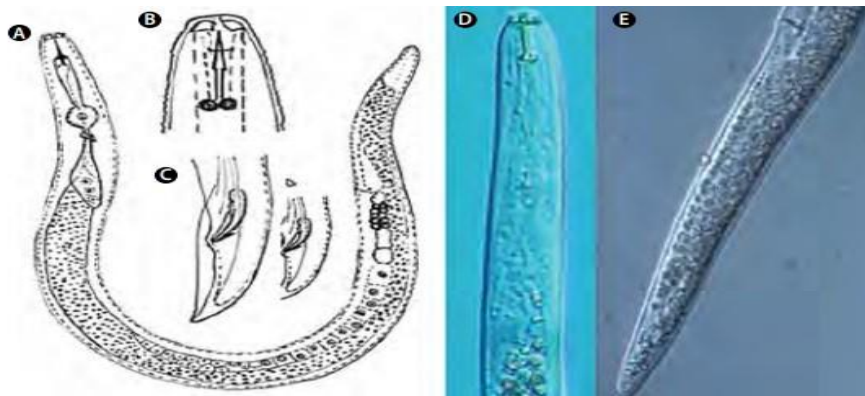


Figure 20. *Pratylenchus* entire body (A), head (B, D), and tail regions (C, E).

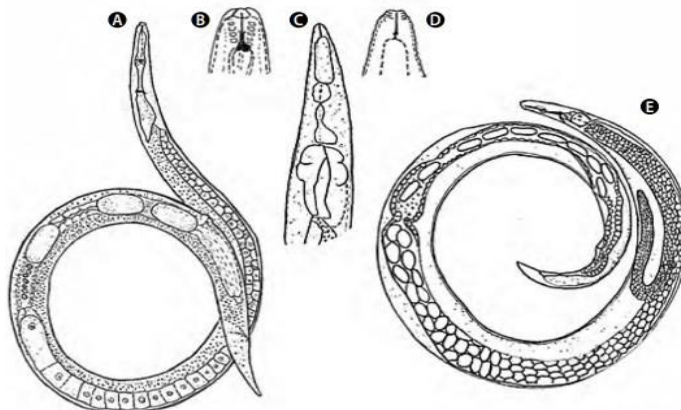


Figure 21. Entire female body of *Ditylenchus* (A), *Ditylenchus* stylet region (B), esophageal region of *Anguina* (C), *Anguina* stylet region (D), and *Anguina* female full body (E).

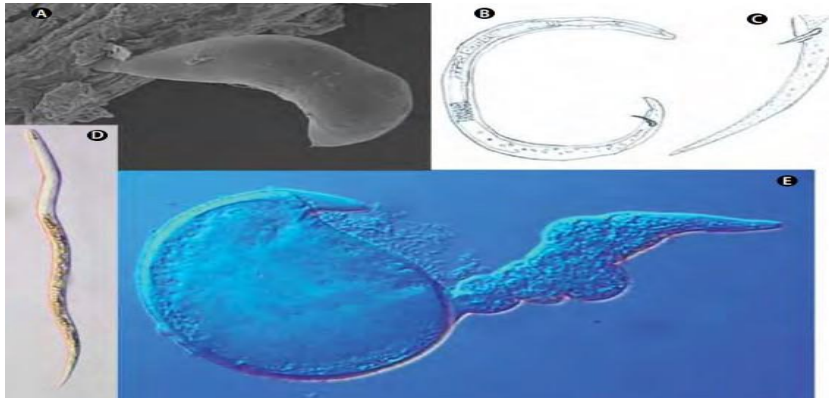


Figure 22. *Rotylenchulus* female protruding from root tissue (A), *Rotylenchulus* male full body (B) and tail region (C), *Tylenchulus* juvenile full body (D) and female full body extracted from root tissue (E).

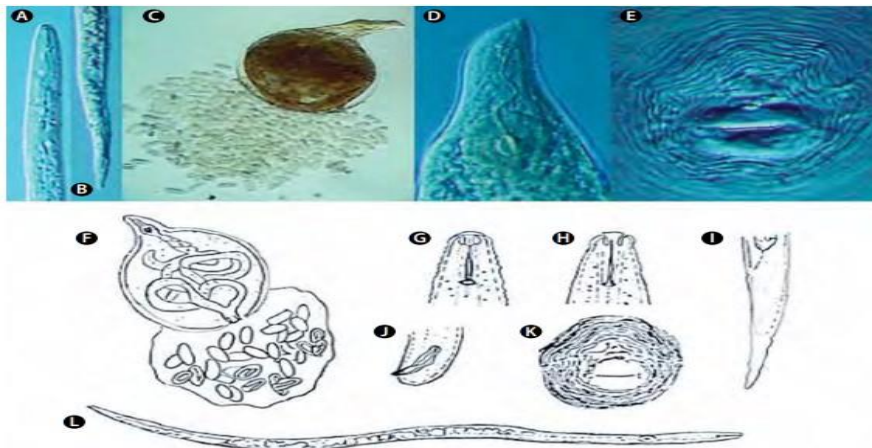


Figure 23. *Meloidogyne* juvenile anterior end (A), juvenile tail (B, I), mature female with eggs (C, F), mature female anterior end (D), female perineal pattern (E), male anterior end (G, H), male tail (J), perineal pattern (K), and juvenile full body (L).

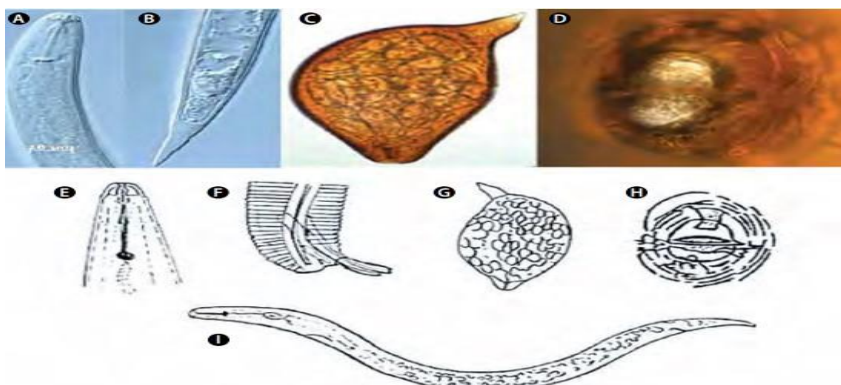


Figure 24. *Heterodera* juvenile anterior end (A), juvenile tail (B), mature cyst with vulval cone (C, G), vulval cone with fenestration (D, H), male anterior end (E), male tail (F), and juvenile full body (I).

Appendix 2: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Initial Stage in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------|
| Treatment | 3 | 336096 | 112032 | 6.505 | 0.0258* |
| Replicate | 2 | 50943 | 25472 | 1.479 | 0.3005 |
| Residuals | 6 | 103340 | 17223 | | |
| Total | 11 | 490379 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 3: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Flowering Stage in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|----------|
| Treatment | 3 | 35826 | 11942 | 5.666 | 0.0348 * |
| Replicate | 2 | 18582 | 9291 | 4.408 | 0.0664. |
| Residuals | 6 | 12646 | 2108 | | |
| Total | 11 | 67054 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 4: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Final Stage in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 250790 | 83597 | 97.790 | 91e-05 *** |
| Replicate | 2 | 1229 | 614 | 0.696 | 0.535 |
| Residuals | 6 | 5291 | 882 | | |
| Total | 11 | 257310 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 5: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Initial Stage in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|--------------|
| Treatment | 3 | 250790 | 83597 | 94.790 | 1.91e-05 *** |
| Replicate | 2 | 1229 | 614 | 0.696 | 0.535 |
| Residuals | 6 | 5291 | 882 | | |
| Total | 11 | 257310 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 6: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Flowering Stage in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|-----------|
| Treatment | 3 | 1155977 | 385326 | 13.331 | 0.00461** |
| Replicate | 2 | 17038 | 8519 | 0.295 | 0.75492 |
| Residuals | 6 | 173425 | 28904 | | |
| Total | 11 | 1346440 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 7: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Final Stage in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|--------------|
| Treatment | 3 | 3198176 | 1066059 | 23.913 | 0.000976 *** |
| Replicate | 2 | 36759 | 18380 | 0.412 | 0.679567 |
| Residuals | 6 | 267488 | 44581 | | |
| Total | 11 | 3502423 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 8: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Initial Stage in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|------------------|----|--------|---------|---------|--------------|
| Treatment | 3 | 27523 | 9174 | 1.013 | 0.4207 |
| Trial | 1 | 49959 | 49959 | 5.519 | 0.0368 |
| Treatment: Trial | 3 | 559363 | 186454 | 20.597 | 5.03e-05 *** |
| Trial: Replicate | 4 | 52172 | 13043 | 1.441 | 0.2801 |
| Residuals | 12 | 108632 | 9053 | | |
| Total | 23 | 797649 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 9: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Flowering Stage in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|------------------|----|---------|---------|---------|--------------|
| Treatment | 3 | 730861 | 243620 | 15.711 | 0.000186 *** |
| Trial | 1 | 36738 | 36738 | 2.369 | 0.149684 |
| Treatment: Trial | 3 | 460941 | 153647 | 9.909 | 0.001440 ** |
| Trial: Replicate | 4 | 35620 | 8905 | 0.574 | 0.686644 |
| Residuals | 12 | 186071 | 15506 | | |
| Total | 23 | 1450231 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 10: ANOVA of the Effect of NPK, TH, and FYM on FLN at the Final Stage in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|------------------|----|---------|---------|---------|--------------|
| Treatment | 3 | 2536477 | 845492 | 37.194 | 2.35e-06 *** |
| Trial | 1 | 735000 | 735000 | 32.334 | 0.000101 *** |
| Treatment: Trial | 3 | 912489 | 304163 | 13.381 | 0.000390 *** |
| Trial: Replicate | 4 | 37988 | 9497 | 0.418 | 0.792802 |
| Residuals | 12 | 272780 | 22732 | | |
| Total | 23 | 4494734 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 11: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Initial Stage in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|--------------|
| Treatment | 3 | 1342095 | 447365 | 73.856 | 3.97e-05 *** |
| Replicate | 2 | 15121 | 7561 | 1.248 | 0.352 |
| Residuals | 6 | 36343 | 6057 | | |
| Total | 11 | 1393559 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 12: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Flowering Stage in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 444144 | 148048 | 38.437 | 0.00026*** |
| Replicate | 2 | 672 | 336 | 0.087 | 0.01760 |
| Residuals | 6 | 23110 | 3852 | | |
| Total | 11 | 467926 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 13: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Final Stage in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------|
| Treatment | 3 | 96051 | 32017 | 4.813 | 0.0488* |
| Replicate | 2 | 24295 | 12147 | 1.826 | 0.2402 |
| Residuals | 6 | 39916 | 6653 | | |
| Total | 11 | 160262 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 14: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Initial Stage in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------|
| Treatment | 3 | 96051 | 32017 | 4.813 | 0.0488* |
| Replicate | 2 | 24295 | 12147 | 1.826 | 0.2402 |
| Residuals | 6 | 39916 | 6653 | | |
| Total | 11 | 160262 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 15: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Flowering Stage in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 437034 | 145678 | 19.239 | 0.00176 ** |
| Replicate | 2 | 40034 | 20017 | 2.644 | 0.15021 |
| Residuals | 6 | 45432 | 7572 | | |
| Total | 11 | 522500 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 16: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Final Stage in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|--------------|
| Treatment | 3 | 2430641 | 810214 | 116.124 | 1.05e-05 *** |
| Replicate | 2 | 295 | 147 | 0.021 | 0.979 |
| Residuals | 6 | 41863 | 6977 | | |
| Total | 11 | 2472799 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 17: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Initial Stage in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|---------|
| Treatment | 3 | 462836 | 154279 | 2.441 | 0.0997. |
| Replicate | 2 | 16446 | 8223 | 0.130 | 0.8789 |
| Trial | 1 | 45763 | 45763 | 0.724 | 0.4067 |
| Residuals | 17 | 1074539 | | | |
| Total | 23 | 1599584 | | | |

Significance codes: 0 '****' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' ' '

Appendix 18: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Flowering Stage in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|--------|
| Treatment | 3 | 78381 | 26127 | 0.496 | 0.690 |
| Replicate | 2 | 16213 | 8106 | 0.154 | 0.859 |
| Trial | 1 | 62424 | 62424 | 1.185 | 0.292 |
| Residuals | 17 | 895832 | 52696 | | |
| Total | 23 | 1052850 | | | |

Significance codes: 0 '****' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' ' '

Appendix 19: ANOVA of the Effect of NPK, TH, and FYM on PPNs at the Final Stage in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|--------------|
| Treatment | 3 | 1727941 | 575980 | 10.954 | 0.000305 *** |
| Replicate | 2 | 11269 | 5634 | 0.107 | 0.898984 |
| Trial | 1 | 83190 | 83190 | 1.582 | 0.225450 |
| Residuals | 17 | 893849 | 52579 | | |
| Total | 23 | 2716249 | | | |

Significance codes: 0 '****' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' ' '

Appendix 20: ANOVA of the Effect of NPK, TH, and FYM on Seed Emergence of Common Beans in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|--------|
| Treatment | 3 | 285.4 | 95.14 | 0.897 | 0.495 |
| Replicate | 2 | 313.5 | 156.77 | 1.478 | 0.301 |
| Residuals | 6 | 636.5 | 106.08 | | |
| Total | 11 | 1235.4 | | | |

Appendix 21: ANOVA of the Effect of NPK, TH, and FYM on Seed Emergence of Common Beans in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|--------|
| Treatment | 3 | 380.7 | 126.9 | 0.496 | 0.699 |
| Replicate | 2 | 501.0 | 250.5 | 0.978 | 0.429 |
| Residuals | 6 | 1536.5 | 256.1 | | |
| Total | 11 | 2418.2 | | | |

Appendix 22: ANOVA of the Effect of NPK, TH, and FYM on Seed Emergence of Common Beans in both trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|--------|
| Trial | 1 | 2 | 2.34 | 0.012 | 0.915 |
| Treatment | 3 | 226 | 75.26 | 0.378 | 0.770 |
| REP | 2 | 40 | 19.79 | 0.099 | 0.906 |
| Residuals | 17 | 3388 | 199.31 | | |
| Total | 23 | 3656 | | | |

Appendix 23: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 14th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 848.25 | 282.75 | 109.33 | <2e-16 *** |
| Replicate | 2 | 8.37 | 4.19 | 1.62 | 0.210 |
| Residuals | 42 | 108.62 | 2.59 | | |
| Total | 47 | 965.25 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 24: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 28th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|------------|
| Treatment | 3 | 2958.75 | 986.25 | 296.67 | <2e-16 *** |
| Replicate | 2 | 11.54 | 5.77 | 1.74 | 0.1886 |
| Residuals | 42 | 139.63 | 3.32 | | |
| Total | 47 | 3109.92 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 25: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 42nd DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 6476.5 | 2158.83 | 352.64 | <2e-16 *** |
| Replicate | 2 | 18 | 9.02 | 1.47 | 0.2407 |
| Residuals | 42 | 257.1 | 6.12 | | |
| Total | 47 | 6751.7 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 26: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 56th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|----------|------------|
| Treatment | 3 | 11202.1 | 3734 | 286.6418 | <2e-16 *** |
| Replicate | 2 | 52.8 | 26.4 | 2.0263 | 0.1445 |
| Residuals | 42 | 547.1 | 13 | | |
| Total | 47 | 11802 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 27: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 14th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 443.58 | 147.86 | 34.78 | 1.848e-11 *** |
| Replicate | 2 | 3.79 | 1.90 | 0.446 | 0.6432 |
| Residuals | 42 | 178.54 | 4.251 | | |
| Total | 47 | 625.92 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 28: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 28th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|---------------|
| Treatment | 3 | 16269.9 | 5423.3 | 58.2436 | 5.137e-15 *** |
| Replicate | 2 | 184.3 | 92.1 | 0.9896 | 0.3802 |
| Residuals | 42 | 3910.8 | 93.1 | | |
| Total | 47 | 20365 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 29: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 42nd DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|--------------|
| Treatment | 3 | 24707 | 8235.8 | 26.8993 | 7.21e-10 *** |
| Replicate | 2 | 1137 | 568.3 | 1.8562 | 0.1689 |
| Residuals | 42 | 12859 | 306.2 | | |
| Total | 47 | 38703 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 30: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 56th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 18843 | 6281.2 | 20.9080 | 1.921e-08 *** |
| Replicate | 2 | 673 | 336.4 | 1.1199 | 0.3359 |
| Residuals | 42 | 12618 | 300.4 | | |
| Total | 47 | 32134 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 31: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 14th DAS in both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|----------|---------------|
| Trial | 1 | 104.17 | 104.17 | 27.8473 | 9.165e-07 *** |
| Treatment | 3 | 1256.67 | 418.89 | 111.9833 | <2e-16 *** |
| Replicate | 2 | 1.58 | 0.79 | 0.2116 | 0.8097 |
| Residuals | 89 | 332.92 | 3.74 | | |
| Total | 95 | 1695.33 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 32: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 28th DAS in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|------------|
| Trial | 1 | 527.3 | 527.3 | 6.4273 | 0.01298 * |
| Treatment | 3 | 16082.3 | 5360.8 | 65.3374 | <2e-16 *** |
| Replicate | 2 | 90.4 | 45.2 | 0.5509 | 0.57840 |
| Residuals | 89 | 7302.2 | 82 | | |
| Total | 95 | 24002.2 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 33: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 42nd DAS in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|----------|------------|
| Trial | 1 | 25026 | 25026 | 134.5693 | <2e-16 *** |
| Treatment | 3 | 28227 | 9409.1 | 50.5945 | <2e-16 *** |
| Replicate | 2 | 676 | 338 | 1.8177 | 0.1684 |
| Residuals | 89 | 16551 | 186 | | |
| Total | 95 | 70481 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 34: ANOVA for Effects of NPK, FYM, and TH on Height of Common Beans at the 56th DAS in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|----------|------------|
| Trial | 1 | 22113 | 22113 | 141.5465 | <2e-16 *** |
| Treatment | 3 | 29481 | 9827.1 | 62.9039 | <2e-16 *** |
| Replicate | 2 | 551 | 275.3 | 1.7624 | 0.1776 |
| Residuals | 89 | 13904 | 156.2 | | |
| Total | 95 | 66049 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 35: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 14th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 45.75 | 15.25 | 9.1337 | 8.982e-05 *** |
| Replicate | 2 | 9.38 | 4.68 | 2.8075 | 0.07172 . |
| Residuals | 42 | 70.13 | 1.67 | | |
| Total | 47 | 125.25 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 36: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 28th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|---------------|
| Treatment | 3 | 2072.06 | 690.69 | 48.1375 | 1.194e-13 *** |
| Replicate | 2 | 286.12 | 143.06 | 9.9708 | 0.0002862 *** |
| Residuals | 42 | 602.62 | 14.35 | | |
| Total | 47 | 2960.81 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 37: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 42nd DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 4844.2 | 1614.75 | 83.265 | < 2.2e-16 *** |
| Replicate | 2 | 514.5 | 257.25 | 13.265 | 3.425e-05 *** |
| Residuals | 42 | 814.5 | 19.39 | | |
| Total | 47 | 6173.2 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 38: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 56th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 6218.1 | 2072.69 | 48.052 | 1.228e-13 *** |
| Replicate | 2 | 947.6 | 473.81 | 10.985 | 0.0001455 *** |
| Residuals | 42 | 1811.6 | 43.13 | | |
| Total | 47 | 8977.3 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 39: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 14th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 45.75 | 15.25 | 12.5588 | 5.364e-06 *** |
| Replicate | 2 | 10.50 | 5.25 | 4.3235 | 0.01962 * |
| Residuals | 42 | 51 | 1.21 | | |
| Total | 47 | 107.25 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 40: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 28th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 2382.0 | 794.0 | 130.202 | < 2.2e-16 *** |
| Replicate | 2 | 148.9 | 74.44 | 12.206 | 6.62e-05 *** |
| Residuals | 42 | 245.6 | 6.10 | | |
| Total | 47 | 2787 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 41: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 42nd DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 31861 | 10620.2 | 95.4399 | <2e-16 *** |
| Replicate | 2 | 251 | 125.4 | 1.1273 | 0.3335 |
| Residuals | 42 | 4647 | 111.3 | | |
| Total | 47 | 36785 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 42: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 56th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 38671 | 12890.2 | 74.7993 | <2e-16 *** |
| Replicate | 2 | 35 | 17.4 | 0.1012 | 0.904 |
| Residuals | 42 | 7238 | 172.3 | | |
| Total | 47 | 45943 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 43: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 14th DAS, Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Trial | 1 | 253.50 | 253.50 | 182.592 | < 2.2e-16 *** |
| Treatment | 3 | 91.50 | 30.50 | 21.969 | 9.791e-11 *** |
| Replicate | 2 | 17.44 | 8.72 | 6.280 | 0.00281 ** |
| Residuals | 89 | 123.56 | 1.39 | | |
| Total | 95 | 486 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 44: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 28th DAS, Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|----------|---------------|
| Trial | 1 | 41.3 | 41.34 | 4.0711 | 0.04664 * |
| Treatment | 3 | 4441 | 1480.34 | 145.7670 | < 2.2e-16 *** |
| Replicate | 2 | 402.9 | 201.47 | 19.8383 | 7.501e-08 *** |
| Residuals | 89 | 903.8 | 10.16 | | |
| Total | 95 | 5789.2 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 45: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 42nd DAS, Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|---------------|
| Trial | 1 | 9362 | 9361.5 | 70.7725 | 6.152e-13 *** |
| Treatment | 3 | 30677 | 10225.8 | 77.3062 | < 2.2e-16 *** |
| Replicate | 2 | 509 | 254.3 | 1.9228 | 0.1522 |
| Residuals | 89 | 11773 | 132.3 | | |
| Total | 95 | 52320 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 46: ANOVA for Effects of NPK, FYM, and TH on Number of Leaves of Common Beans at the 56th DAS, Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Trial | 1 | 16695 | 16695.4 | 89.8894 | 3.791e-15 *** |
| Treatment | 3 | 37760 | 12586.8 | 67.7682 | < 2.2e-16 *** |
| Replicate | 2 | 630 | 315.1 | 1.6965 | 0.1892 |
| Residuals | 89 | 16530 | 185.7 | | |
| Total | 95 | 71616 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 47: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 14th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 5.08 | 1.69 | 9.1337 | 8.982e-05 *** |
| Replicate | 2 | 1.04 | 0.52 | 2.8075 | 0.07172 . |
| Residuals | 42 | 7.79 | 0.19 | | |
| Total | 47 | 13.92 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 48: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 28th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 230.23 | 76.74 | 48.1375 | 2.45e-13 *** |
| Replicate | 2 | 31.79 | 15.90 | 9.9708 | 0.0002862 *** |
| Residuals | 42 | 66.96 | 1.59 | | |
| Total | 47 | 328.98 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 49: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 42nd DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 538.25 | 179.42 | 83.265 | < 2e-16 *** |
| Replicate | 2 | 57.17 | 28.58 | 13.265 | 3.425e-05 *** |
| Residuals | 42 | 90.50 | 2.15 | | |
| Total | 47 | 685.92 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 50: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 56th DAS in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 692.7 | 230.91 | 50.987 | 4.683e-14 *** |
| Replicate | 2 | 113.37 | 56.69 | 12.517 | 5.445e-05 *** |
| Residuals | 42 | 190.21 | 4.53 | | |
| Total | 47 | 996.31 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 51: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 14th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 5.08 | 1.69 | 12.56 | 5.364e-06 *** |
| Replicate | 2 | 1.17 | 0.58 | 4.32 | 0.01962 * |
| Residuals | 42 | 5.67 | 0.13 | | |
| Total | 47 | 11.92 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 52: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 28th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Treatment | 3 | 264.67 | 88.22 | 130.202 | < 2e-16 *** |
| Replicate | 2 | 16.54 | 8.27 | 12.206 | 6.621e-05 *** |
| Residuals | 42 | 28.46 | 0.68 | | |
| Total | 47 | 309.67 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 53: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 42nd DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 3540.1 | 1180.03 | 95.4399 | <2e-16 *** |
| Replicate | 2 | 27.9 | 13.94 | 1.1273 | 0.3335 |
| Residuals | 42 | 519.3 | 12.36 | | |
| Total | 47 | 4087.2 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 54: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 56th DAS in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|------------|
| Treatment | 3 | 4296.7 | 1432.24 | 74.7993 | <2e-16 *** |
| Replicate | 2 | 3.9 | 1.94 | 0.1012 | 0.904 |
| Residuals | 42 | 804.2 | 19.15 | | |
| Total | 47 | 5104.8 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 55: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 14th DAS in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Trial | 1 | 28.167 | 28.167 | 182.59 | < 2.2e-16 *** |
| Treatment | 3 | 10.167 | 3.3889 | 21.969 | 9.791e-11 *** |
| Replicate | 2 | 1.937 | 0.969 | 6.280 | 0.00218 ** |
| Residuals | 89 | 13.729 | 0.15 | | |
| Total | 95 | 54 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 56: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 28th DAS in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|----------|---------------|
| Trial | 1 | 4.59 | 4.59 | 4.0711 | 0.04664 * |
| Treatment | 3 | 493.45 | 164.48 | 145.7670 | < 2.2e-16 *** |
| Replicate | 2 | 44.77 | 22.39 | 19.8383 | 7.501e-08 *** |
| Residuals | 89 | 100.43 | 1.10 | | |
| Total | 95 | 643.24 | 1.13 | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 57: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 42nd DAS in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Trial | 1 | 1040.2 | 1040.17 | 70.7725 | 6.152e-13 *** |
| Treatment | 3 | 3408.6 | 1136.19 | 77.3062 | < 2.2e-16 *** |
| Replicate | 2 | 56.5 | 28.26 | 1.9228 | 0.1522 |
| Residuals | 89 | 1308.1 | 14.70 | | |
| Total | 95 | 5813.3 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 58: ANOVA for Effects of NPK, FYM, and TH on Number of Branches of Common Beans at the 56th DAS in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr (>F) |
|-----------|----|--------|---------|---------|---------------|
| Trial | 1 | 1837.5 | 1837.5 | 89.6372 | 4.039e-15*** |
| Treatment | 3 | 4201.9 | 1400.63 | 68.3255 | < 2.2e-16 *** |
| Replicate | 2 | 74.8 | 37.41 | 1.8248 | 0.1672 |
| Residuals | 89 | 1824.4 | 20.50 | | |
| Total | 95 | 7938.6 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 59: ANOVA for Effects of NPK, FYM, and TH on the number of pods of Common Beans in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|-----|--------|---------|----------|------------|
| Treatment | 3 | 6061.4 | 2020.5 | 219.8488 | <2e-16 *** |
| Replicate | 2 | 6.6 | 3.32 | 0.3615 | 0.6972 |
| Residuals | 114 | 1047 | 9.19 | | |
| Total | 119 | 7115.7 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 60: ANOVA for Effects of NPK, FYM, and TH on Number of Pods of Common Beans in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|-----|--------|---------|----------|------------|
| Treatment | 3 | 6594.2 | 2198.05 | 241.0872 | <2e-16 *** |
| Replicate | 2 | 55.8 | 27.9 | 3.0601 | 0.05075 . |
| Residuals | 114 | 1039.4 | 9.12 | | |
| Total | 119 | 7689.3 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 61: ANOVA for Effects of NPK, FYM, and TH on Number of Pods of Common Beans in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|-----|---------|---------|----------|------------|
| Trial | 1 | 3 | 3 | 0.3342 | 0.5638 |
| Treatment | 3 | 12647.5 | 4215.8 | 463.7822 | <2e-16 *** |
| Replicate | 2 | 39.5 | 19.7 | 2.1713 | 0.1163 |
| Residuals | 233 | 2118 | 9.1 | | |
| Total | 239 | 14808.1 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 62: ANOVA for Effects of NPK, FYM, and TH on Number of Seeds of Common beans in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|-----|--------|---------|----------|---------------|
| Treatment | 3 | 359597 | 119866 | 145.9167 | < 2.2e-16 *** |
| Replicate | 2 | 10244 | 5122 | 6.2353 | 0.002693 ** |
| Residuals | 114 | 93647 | 821 | | |
| Total | 119 | 463489 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 63: ANOVA for Effects of NPK, FYM, and TH on Number of Seeds of Common beans in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|-----|--------|---------|----------|------------|
| Treatment | 3 | 282889 | 94296 | 216.8836 | <2e-16 *** |
| Replicate | 2 | 969 | 485 | 1.1145 | 0.3316 |
| Residuals | 114 | 49565 | 435 | | |
| Total | 119 | 333422 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 64: ANOVA for Effects of NPK, FYM, and TH on Number of Seeds of Common beans in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|-----|--------|---------|----------|---------------|
| Trial | 1 | 1728 | 1728 | 2.6680 | 0.10581 |
| Treatment | 3 | 639170 | 213057 | 328.9388 | < 2.2e-16 *** |
| Replicate | 2 | 6825 | 3412 | 5.2684 | 0.005784 ** |
| Residuals | 233 | 150916 | 648 | | |
| Total | 239 | 798639 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 65: ANOVA for Effects of NPK, FYM, and TH on Yield of Common Beans in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|----------|---------|---------|------------|
| Treatment | 3 | 23220966 | 7740322 | 15.70 | 0.00302 ** |
| Replicate | 2 | 1035469 | 517735 | 1.05 | 0.40644 |
| Residuals | 6 | 2958445 | 493074 | | |
| Total | 11 | 27214880 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 66: ANOVA for Effects of NPK, FYM, and TH on Yield of Common Beans in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|----------|---------|---------|------------|
| Treatment | 3 | 27427082 | 9142361 | 11.16 | 0.00723 ** |
| Replicate | 2 | 2900776 | 1450388 | 1.77 | 0.24876 |
| Residuals | 6 | 4916330 | 819388 | | |
| Total | 11 | 35244188 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 67: ANOVA for Effects of NPK, FYM, and TH on Yield of Common Beans in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|----------|----------|---------|--------------|
| Trial | 1 | 1731616 | 1731616 | 3.55 | 0.0768 . |
| Treatment | 3 | 50556934 | 16852311 | 34.55 | 1.87e-07 *** |
| Replicate | 2 | 3609499 | 1804750 | 3.70 | 0.0464 * |
| Residuals | 17 | 8292635 | 487802 | | |
| Total | 23 | 64190684 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 68: ANOVA for Effects of NPK, FYM, and TH on Biomass of Common beans in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|--------------|
| Treatment | 3 | 1989632 | 663211 | 64.529 | 5.88e-05 *** |
| Replicate | 2 | 16097 | 8049 | 0.783 | 0.499 |
| Residuals | 6 | 61666 | 10278 | | |
| Total | 11 | 2067395 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 69: ANOVA for Effects of NPK, FYM, and TH on Biomass of Common Beans in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|--------------|
| Treatment | 3 | 1637175 | 545725 | 122.991 | 8.91e-06 *** |
| Replicate | 2 | 7873 | 3937 | 0.887 | 0.46 |
| Residuals | 6 | 26623 | 4437 | | |
| Total | 11 | 1671671 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 70: ANOVA for Effects of NPK, FYM, and TH on Biomass of Common beans in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|--------------|
| Trial | 1 | 2136 | 2136 | 0.323 | 0.577 |
| Treatment | 3 | 3603740 | 1201247 | 181.431 | 4.17e-13 *** |
| Replicate | 2 | 22770 | 11385 | 1.720 | 0.209 |
| Residuals | 17 | 112556 | 6621 | | |
| Totals | 23 | 3741202 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 71: ANOVA for Effect of NPK, FYM, and TH on Number of Nodules in Common Beans in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|---------------|
| Treatment | 3 | 8290.9 | 2763.63 | 42.8003 | 5.877e-11 *** |
| Replicate | 2 | 160.2 | 80.11 | 1.2407 | 0.3036 |
| Residuals | 30 | 1937.1 | 64.57 | | |
| Total | 35 | 10388.2 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 72: ANOVA for Effect of NPK, FYM, and TH on Number of Nodules in Common Beans in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|---------|---------------|
| Treatment | 3 | 15809.9 | 5270 | 240.292 | < 2.2e-16 *** |
| Replicate | 2 | 693.2 | 346.6 | 15.803 | 2.053e-05 *** |
| Residuals | 30 | 657.9 | 21.9 | | |
| Total | 35 | 17161.0 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 73: ANOVA for Effect of NPK, Fym, and Th on Number of Nodules in Common Beans in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|---------|---------|----------|---------------|
| Trial | 1 | 2913.4 | 2913.4 | 50.1713 | 1.237e-09 *** |
| Treatment | 3 | 23138.4 | 7712.8 | 132.8217 | < 2.2e-16 *** |
| Replicate | 2 | 636.4 | 318.2 | 5.4794 | 0.006323 ** |
| Residuals | 65 | 3774.5 | 58.1 | | |
| Total | 71 | 30462.6 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 74: ANOVA for Effect of NPK, FYM, and TH on PPNs in Roots of Common Beans in Trial One

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|----------|
| Treatment | 3 | 3721 | 1240.4 | 4.776 | 0.00496* |
| Replicate | 2 | 133 | 66.3 | 0.255 | 0.7827 |
| Residuals | 6 | 1558 | 259.7 | | |
| Total | 11 | 5412 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 75: ANOVA for Effect of NPK, FYM, and TH on PPNs in Roots of Common Beans in Trial Two

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|-----------|----|--------|---------|---------|--------------|
| Treatment | 3 | 7388 | 2462.8 | 36.575 | 0.000299 *** |
| Replicate | 2 | 248 | 124.0 | 1.842 | 0.237904 |
| Residuals | 6 | 404 | 67.3 | | |
| Total | 11 | 8040 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 76: ANOVA for Effect of NPK, FYM, and TH on PPNs in Roots of Common Beans in Both Trials

| | DF | Sum Sq | Mean Sq | F value | Pr(>F) |
|------------------|----|--------|---------|---------|--------------|
| Treatment | 3 | 9488 | 3163 | 19.341 | 6.86e-05 *** |
| Trial | 1 | 1610 | 1610 | 9.848 | 0.00856 ** |
| Trial: Replicate | 4 | 381 | 95 | 0.582 | 0.68161 |
| Treatment: Trial | 3 | 1622 | 541 | 3.306 | 0.05741 . |
| Residuals | 12 | 1962 | 164 | | |
| Total | 23 | 15063 | | | |

Significance codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Appendix 77: Pictorial Presentation of Field Experiment and Laboratory Work



Appendix 78: Chuka University Ethics Approval letter



CHUKA UNIVERSITY INSTITUTIONAL ETHICS REVIEW COMMITTEE

Telephones: 020-2310512/18

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P. O. Box 109-60400, Chuka

Website: www.chuka.ac.ke

19th March, 2024

REF: CUIERC/ NACOSTI/490
TO: Margaret Wairimu Muraki

RE: Soil Nematode Communities under Different Fertility Management Practices, Biocontrol, Growth and Yield of Common Beans (var.Mwitmania) in Tharaka Nithi, Kenya

This is to inform you that *Chuka University IERC* has reviewed and approved your above research proposal. Your application approval number is *NACOSTI/NBC/AC-0812*. The approval period is 19th March, 2024 – 19th March, 2025.

This approval is subject to compliance with the following requirements;


- i. Only approved documents including (informed consents, study instruments, MTA) will be used
- ii. All changes including (amendments, deviations, and violations) are submitted for review and approval by *Chuka University IERC*.
- iii. Death and life threatening problems and serious adverse events or unexpected adverse events whether related or unrelated to the study must be reported to *Chuka University IERC* within 72 hours of notification
- iv. Any changes, anticipated or otherwise that may increase the risks or affected safety or welfare of study participants and others or affect the integrity of the research must be reported to *Chuka University IERC* within 72 hours
- v. Clearance for export of biological specimens must be obtained from relevant institutions.
- vi. Submission of a request for renewal of approval at least 60 days prior to expiry of the approval period. Attach a comprehensive progress report to support the renewal.
- vii. Submission of an executive summary report within 90 days upon completion of the study to *Chuka University IERC*.

Prior to commencing your study, you will be expected to obtain a research license from National Commission for Science, Technology and Innovation (NACOSTI) <https://oris.nacosti.go.ke> and also obtain other clearances needed.

Yours sincerely

Dr. Benjamin Kanga
SECRETARY

Appendix 79: NARCOSTI Research Permit




NATIONAL COMMISSION FOR SCIENCE, TECHNOLOGY & INNOVATION.

RESEARCH LICENSE

Ref No: **197878**


Date of Issue: **06/May/2024**




This is to Certify that Miss. Margaret Wairimu Muraki of Chuka University, has been licensed to conduct research as per the provision of the Science, Technology and Innovation Act, 2013 (Rev.2014) in Tharaka-Nithi on the topic: SOIL NEMATODE COMMUNITIES UNDER DIFFERENT FERTILITY MANAGEMENT PRACTICES, BIOCONTROL, GROWTH AND YIELD OF COMMON BEANS (var. Mwittemania) IN THARAKA NITHI, KENYA for the period ending : 06/May/2025.

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197878
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